THE U.S. DEPARTMENT OF HEALTH AND HUMAN SERVICES PUBLIC HEALTH SERVICE CENTERS FOR DISEASE CONTROL AND PREVENTION NATIONAL INSTITUTE FOR OCCUPATIONAL SAFETY AND HEALTH

convenes the

NINETEENTH MEETING

ADVISORY BOARD ON

RADIATION AND WORKER HEALTH

The verbatim transcript of the Meeting of the Advisory Board on Radiation and Worker
Health held at The Westin Casuarina, 160 East Flamingo Road, Las
Vegas, Nevada, on December 9 and 10, 2003.

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COURT REPORTER'S CERTIFICATION

TRANSCRIPT LEGEND

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(By Group, in Alphabetical Order)

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(in order of appearance)

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Mr. Jeff Kotsch, DOL

Mr. Stu Hinnefeld, NIOSH

Mr. David Allen, NIOSH

Dr. James Melius, Workgroup Chair

Dr. John Mauro, SC&A

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JEFF KOTSCH
JOHN MAURO
RICHARD MILLER
DAVID NAIMON

LOUISE S. PRESLEY KNUTE RINGIN CHUCK ROESSLER BRIAN THOMAS RICHARD TOOHEY

PROCEEDINGS

2 Error!

3 (8:30 a.m.)

REGISTRATION AND WELCOME

5 DR. ZIEMER: Good morning. I'd like to call to order the
19th meeting of the Advisory Board on Radiation and
Worker Health. I'm Paul Ziemer, Chairman of the Board,
and the other members of the Board are all present and
you see their names on the placards before them.
This is a slightly different venue than we're used to. If
we do well today, I understand that we may become

permanent replacements for Siegfried and Roy, but we'll see how it goes. If you're here expecting to hear David Brenner, you're here too early today. Besides, he charges \$50 a person and this show is free. And they say you get what you pay for.

7 I'd like to welcome not only staff from several Federal
8 agencies, but other members of the public who may be
9 here this morning. We'd like to remind all of you --

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Board members, Federal staff, contractor staff and
      members of the public -- to please register your
      attendance in the book that's out at the entrance.
      Also members of the public, if you wish to make public
      comment during one of the public comment periods, the
      first of which will be about mid-afternoon -- 2:45 --
      to please sign up at that registration book so that we
      have some idea of how many will be presenting at that
      time.
O There are copies of a number of items on the table over here
      on my left near the rear. Some are presentations that
      will occur today. There's copies of the agenda.
      are copies of past Board minutes and other related
      items, so please avail yourself of that information, as
      well.
6 Finally, we do welcome you to Las Vegas. This seems like an
      appropriate place to talk about probabilities and
      risks, although I'm sure the view here gets distorted
      because people seem to think they can beat the
      probabilities.
                      In any event, let me turn the mike over
      briefly to Larry Elliott to add a few comments.
2 MR. ELLIOTT: On behalf of the Director of NIOSH, John
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Howard, and the Secretary of Health and Human Services,
      Secretary Thompson, I'd like to welcome the Board to
      Las Vegas. I'd like to welcome the public to this
      meeting. It is a public meeting.
5 We would like to apologize for the particular forum that we
      are presented in here today. We typically use a
      different forum, a more level playing field, if you
      will, where the Board is on the same level with the
      audience. Unfortunately, in this hotel, the room that
      we had contracted was not available for us.
      got a building inspection permit and there's an egress
      problem, so this was the only other space that was
      available to us. And because we have announced this
      public meeting in the Federal Register, we must hold it
      here at this address. And so we'll make do here today;
      I beg all your indulgences.
7 We have a busy agenda, and I look forward to a productive
      meeting.
                Thank you.
9 DR. ZIEMER:
              Thank you, Larry. The minutes for meeting 18,
      which was the meeting held in St. Louis, Missouri in
      October, were provided to the Board in their books,
      which many of them only got this morning. And the
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minutes are fairly lengthy, so without objection, I'm
      going to defer action on the minutes until our meeting
      tomorrow.
4 MS. MUNN:
            Thank you.
5 DR. ZIEMER:
              I do ask the Board members in the meantime to
      spend their evening in their rooms reading the Board
      minutes. But more seriously and particularly, look at
      those areas in the minutes which summarize discussions
      that you might have contributed to to make sure that we
      have accurately rendered your thoughts. For example,
      I'm going to ask Dr. Roessler to check her thoughts --
      the rendering of her thoughts because I can't figure
      them out in the minutes --
4 DR. ROESSLER: I'll see if I can.
5 DR. ZIEMER: -- and I'm hopeful that she can, and there may
      be others like that, as well. I'm not picking on her,
      but I don't think the minutes maybe fully reflected
      what was being said. But particularly check those
      areas where there is sort of a summary of what your
      views or thoughts were on particular issues.
1 And if you have other changes that you would like to make,
      we'd like to have those by tomorrow. And Ray, our
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court recorder who also prepares the summary minutes for us, will be able to incorporate those changes before we issue the final copy. So be prepared 3 tomorrow to identify any substantive changes and then we'll also get from you -- and you can turn them in on your own -- any grammatical changes that you might note.

8 Are there any questions on those minutes other than the remarks I've just made?

(No responses)

1 DR. ZIEMER: Okay, thank you very much. Then we will defer action on those until tomorrow.

PROGRAM STATUS REPORT

4 We'd like to proceed then with the next item on the agenda, which is the program status report. Chris Ellison's going to present that for us today. I should point out, particularly for those in the audience, that because of the setup and the venue here, it may be a little awkward for you. The screen is over here on my right and about in the middle of the room, so those near the front may have to do a bit of twisting and turning to see the screen.

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1 Chris, glad to have you here today. Please proceed.
2 MS. ELLISON: Good morning. Can everyone hear me okay?
3 UNIDENTIFIED:
               No.
4 MS. ELLISON: Okay, before I proceed --
5 DR. ZIEMER:
              No.
6 MS. ELLISON: No?
7 DR. ZIEMER: Maybe a little closer, or they'll turn it up a
      little, perhaps.
9 MS. ELLISON: Better?
O DR. ZIEMER:
              Yes.
1 MS. ELLISON:
               Much better. This morning I'll be presenting
      the current program overview, and it's very similar to
      a lot of the format that you've seen before when Dave
3
      Sundin has been presenting this -- similar format.
      then when I get done with this presentation, I also
      would like to show you a web page that we're working
           It's a different look at a lot of the program
      stats that we currently have.
9 MR. ELLIOTT: Chris, if you put that on the right side, when
      you talk to the -- the screen, it'll pick your voice up
      better.
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2 MS. ELLISON: Better? Now as you know, we have started

receiving cases from the Department of Labor back in October of 2001, and this slide here shows the number of cases that we've received currently, based on each year. And at the current time, we're just under 15,000 cases that we've received from the Department of Labor. And if things go at the current rate, it does appear that we'll probably hit 15,000 probably by the end of December. 9 Now with those 15,000 cases that we have received, the number of cases that are considered Atomic Weapons Employer, or AWE, cases are just over 2,000, which is roughly still 14 percent, which is what it's been running now for quite some time. And then the number of cases that are non-AWE employees, which are the Department of Energy employees, is just a little more than 17,500 (sic). 7 Now if you do your math real quick with those two numbers -when I was putting together the presentation I added those numbers and I though that doesn't quite add up to the 14,895 that we're showing, so I did a little bit of investigative work, wondering why that was occurring, 2 and I was told that more than likely the Department of

Energy has sent us some beryllium cases that we should not have had. 3 Which leads me into the last number on this slide, which is the number of cases that we currently have in process or in-house that we're working on the dose reconstructions, which is just a little over 13,500. And those numbers are the active cases. It does not include cases that we have returned to the Department of Labor because we've completed the dose reconstructions. And there are some cases that are sent to us by mistake, which would include the beryllium cases, and sometimes we receive duplicate So that 13,563 is actually what we have inhouse and we're working on the dose reconstructions right now. 6 This is a trend chart, and I know you've seen it before, also. And what I've done is I've updated the fourth quar-- or the first quarter of fiscal year 2004. based on this, you can see that the numbers are still continuing to decline in the number of cases that we receive from the Department of Labor. However, it does 2 average out -- this is based at the end of November.

It is still averaging out just slightly over 200 a month. 3 One of the first things that happens to the cases when we receive them from the Department of Labor, we do have to scan everything in that we receive and put it into the electronic database that we have. Once that is completed, the next thing that we do with the case is we do send a request to the Department of Energy for the exposure monitoring information. And this here shows the number of requests that we have currently sent to the Department of Energy, and that's just over 14,000 requests. That does represent roughly 12,700 claims or cases that we have. And keep in mind that a lot of these cases, if someone has worked at multiple sites, we do have to send out more than one request to cover all of the sites that they have worked at. 7 Thus it gets to the next item up there, the responses we've And that's why that's a little bit higher received. than the number of total requests that we've sent out, which the responses now at almost 22,000. accounts for the multiple sites that individuals have 2 worked at, and then also sometimes the Department of

2

Energy sends us more than request -- or response to our request. They'll send it in partial pieces.

The last item on this slide shows the number of outstanding requests that we have. We do ask that the Department of Energy try to fill our requests within 60 days, and we have been working with them in trying to get those requests in a timely manner. And here are some points and some days that the numbers are showing of the number of cases that are currently at those 60, 90, 120 and 150 days.

But I want to draw your attention to this next slide. I

know this next slide -- you normally see it with some percentages off to the side, and I wanted to pull the percentages off to kind of make the numbers pop out and stand out a little bit more. And I do want to note a few things about this. We've been saying that we're continuing to see a little bit better response rate in getting the information to us in a more timely manner from the Department of Energy. And a couple of the sites I want to note that are on here have increased their percentage response rate from October. Savannah River Site in October they were showing a 78 percent

response rate; they are now showing an 86 percent response rate, so it has gone up a few percentages for Savannah River. 4 Richland has also increased their response rate. It went 5 from a 94 percent to a 98. Nevada Test Site, two percent increase; it went from a 97 to a 99 percent. And the last one that showed a fairly significant increase was the Idaho Operations Office, and they went from a 57 percent response rate to -- or a 50 percent response rate, excuse me, to a 57, so the continued meetings that we do have with the Department of Energy appear to be working. We must keep in mind also that, you know, the Department of Energy had to get their programs up and running in order to provide us these responses, so it does show that they are responding to the requests. 7 Now the telephone interviews. As you know, the Act did not require that we conduct individual telephone interviews with the claimants, but we felt we wanted to build it into the process because it gave the -- gives the claimant the opportunity to provide us with any 2 additional information that they may be aware of that

will help us in doing the dose reconstruction for their case. So currently our contractor, Oak Ridge Associated Universities, is working on the telephone interviews for us. And if you look, they've conducted at least one telephone interview for almost 9,000 And with -- once the telephone interview has been conducted, we do send a summary report to the claimant, and we ask them to review the summary report and provide us with any updates or corrections that they may see. And I'm showing that not quite 11,500 of those summary reports have been sent out. current capacity that ORAU is showing in conducting the interviews is still maintained at about 200 to 300 a week.

5 Now the information that everyone's been waiting on, the dose reconstructions. In previous presentations there was one category called cases initiated. We now have the ability to break that into two different parts, and the first one is cases staged for dose reconstruction, which is roughly 2,700. What this means is these are cases who have all their telephone interviews completed. We have received dose information to do the

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dose reconstruction. We've sent them a letter telling
      them that we've received all that information and, if
      applicable, a site profile has been completed for their
      site. Those cases are ready to be assigned.
5 Therefore we have the next number is the number of DRs
      assigned. These are one -- cases that are currently
      assigned to a health physicist and we are working on
      the dose reconstructions, and that is at 631.
9 The number of draft reports that are sent to a claimant are
      just about at 250, and those are ones where we've sent
      the dose -- draft dose reconstruction report to the
      claimant and asked them to review it. We will conduct
      a close-out interview with the claimant to explain the
      draft dose reconstruction report to them. And we also
      send at that time the OCAS-1 form and asking them to
      sign that and return that to us.
7 And the last figure for the dose reconstruction statistics
      are the final ones sent to DOL. We also do send a copy
      to DOE, and those are just at over 1,000. Actually I
      called this morning. That number's gone up slightly.
      It's at 1,045. I wanted to see -- I did these stats
2
      Friday morning. I wanted to see how much it changed
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over the weekend at the end of that day.
2 The phone calls and the e-mails that we receive -- one thing
      that I reported last time at the October meeting was
3
      that we were going to be sending out an activity
               And if you look at the numbers -- there are
      large numbers that are up here. OCAS phone calls are
      almost at 26,000. ORAU phone calls are right around
      53,000. And there's a little bit of a reason for that
      difference. The phone calls that OCAS primarily takes
      are claimant calls, and ORAU contractor -- they do set
      up telephone interviews, and each time they attempt to
      contact a claimant, that is logged into our database.
      So that can account for why their numbers are slightly
      higher.
5 However, since we did send the activity report out, I know
      that our public health advisors within OCAS have
      commented that the phones haven't been ringing quite as
      much, and I did look and they're down from -- let's
      see, October to November, the phone calls are down
      about 600 each month. They're a slight decline there.
1 And the e-mails are pretty constant, still coming in and
      total within the system is just over 2,700 (sic).
```

1 And now the recent accomplishments that we've had with the program since the last meeting. In November we did appoint 36 additional physicians to the DOE physician panels, which brings the total to 159. Also, the residual contamination final report was released, and it is available on our web site. And I did mention the third item, we hit a milestone -- I believe it was last week -- with the 1,000 DRs completed and sent back to the Department of Labor.

0 Now the last item on here -- these are also available on the web site -- are the various site profiles. And we have been publishing them and putting them on the web site once the items are available. From the time of the October meeting until now, we've placed several new documents on the web site, and I did try to remember to send an e-mail to the Advisory Board to let you know when this has been done. We've added an AWE and a DOE site-wide documents, I believe there are two for the DOE facilities. Also the Hanford site profile is now complete. I just posted the introduction on the web There is now a site profile for the Huntington site. Pilot Plant and with INEEL, Portsmouth and X-10, we

- posted part of the site profile. It was -- the site
- description has been posted, and the other pieces are
- now pending. And then we have Y-12 with the site
- description and the dosimetry.
- 5 That's all I have basically for the stats. Are there any
- 6 questions before I go on?
- 7 DR. ZIEMER: Thank you, Chris. Let me begin with a couple
- 8 of points for clarification.
- 9 MS. ELLISON: Sure.
- O DR. ZIEMER: The slide number -- I think it's slide six, you
- have interview summary reports sent to claimants,
- 2 11,499.
- 3 MS. ELLISON: Uh-huh.
- 4 DR. ZIEMER: And you also listed cases for which one or more
- 5 interview is completed, 8,954. Clarify the difference
- in those two numbers.
- 7 MS. ELLISON: The 8,954 represents the number of telephone
- 8 interviews for at -- at least of which one interview is
- 9 conducted.
- 0 DR. ZIEMER: One or more.
- 1 MS. ELLISON: At least one.
- 2 DR. ZIEMER: Right.

```
1 MS. ELLISON: Right, and with a lot of the cases there are
      multiple survivors, each survivor or applicant has the
      opportunity to participate --
4 DR. ZIEMER:
              So there are multiple summaries --
5 MS. ELLISON:
               Right.
6 DR. ZIEMER:
              -- in some cases then.
7 MS. ELLISON: That 8,954 represents per case --
8 DR. ZIEMER:
              Gotcha.
9 MS. ELLISON: -- that's the total number of cases.
O DR. ZIEMER:
              Okay.
1 MS. ELLISON:
               Where the summary is total numbers of summary,
      not...
3 DR. ZIEMER: And then on the next slide where you give
      number of draft reports sent to claimants, I assume
      that's just the number that are currently out there --
6 MS. ELLISON: Right, that we're waiting --
7 DR. ZIEMER:
             -- that last column, final draft reports, at
      one time -- or the final -- the final reports, not
      draft --
0 MS. ELLISON: Uh-huh.
1 DR. ZIEMER: -- at one time were drafts, so that this 249
      drafts is just what's currently --
```

- 1 MS. ELLISON: Right, that's currently what we're waiting on that OCAS-1 to come back. There's 249 out there.
- 3 DR. ZIEMER: Thank you.
- 4 MS. ELLISON: Uh-huh. One more thing -- any other questions
- on the slide presentation?
- 6 DR. ZIEMER: There's one question here, start with Bob, then
- we'll come down.
- 8 MR. PRESLEY: Do you think --
- 9 DR. ZIEMER: Use the mike, Robert.
- 0 MR. PRESLEY: Robert Presley. Chris, do you have any reason
- why that Savannah River and Idaho and Los Alamos -- has
- 2 DOE given you any reason why that they've got so many
- that are better than 150 days?
- 4 MS. ELLISON: I know that we have been in contact with them
- and we are working with them and they are setting up
- 6 their systems. A lot of those -- those three that you
- named off, I didn't provide their percentages because
- 8 they're maintaining the constant level from the last
- time, so -- I do know that we are working with them in
- trying to get those, and have identified specifically
- 1 which of the cases that we're waiting on so they are
- aware of that and they are working on them.

```
1 MR. PRESLEY:
               Thank you.
2 MS. ELLISON:
               Uh-huh.
3 DR. ZIEMER:
             Okay, Jim?
4 DR. MELIUS: Yeah, just to follow up on that, the -- those -
      - you have a large number of cases that are over 150
      days for those -- I think those three sites that were
      just mentioned. Are those sites -- I take it those go
      -- are any of those ever getting cleared or is that
      just sort of a steady, you know, number --
0 MS. ELLISON: No, we are receiving --
1 DR. MELIUS:
             -- constant -- but -- but are we receiving --
      are -- is there a group out there that they're not
      finding records on or are they just -- is that just a
3
      question of the flow through the system as things come
      back from -- 'cause that's a long time for a claimant
      to wait and --
7 MS. ELLISON:
               Right. In the monthly reports we do specify
      which claims or cases that we are waiting on the
      information, and we do require the Department of Energy
      to send us a response saying no, they cannot find any
      information. So in those cases they have not indicated
2
      that they cannot find anything.
```

I think Larry has some additional comments and 1 DR. ZIEMER: 3 DR. MELIUS: Yeah, let me ask one additional point 'cause then Larry can answer them both. Is the Iowa -- the Iowa site, which isn't listed there, but if -- as I recall from a couple of meetings ago, there was a major problem in getting records from that site and has that been resolved? Yes, it has. I'll let Larry answer that. 9 MS. ELLISON: O MR. ELLIOTT: To your first question, let me respond that some of those numbers that you see there are static. They've been there from the start and we're working on those individual cases with that respective site to 3 understand better what is going on. And in some cases, yes, they are having trouble even verifying that the person actually, in their records, worked at the site, perhaps, because they might have been verified by Labor as having worked at the site by IRS records or some other sort of record system. The majority of those -those numbers that you saw, though, change. it's a small group that we see as static, and we're 2 working on all of the numbers.

```
1 The Idaho site had to index -- had to scan all of their
      boxes of records -- thousands of boxes of records, and
      then index through that scanning effort so that they
      could retrieve a person's history of dose. And that
      took a while to do and that has been completed.
      fact, our ORAU team has aided the Idaho site by
      providing some further contract support to people who
      are working on that indexing effort who are now
      searching for those records for those individuals.
      we've accommodated that through that particular site.
1 Now to your second question, the Iowa plant, yes, that -- we
      have I think received five or six boxes of records that
      were held by the Department of Defense, and that's been
3
      very critical. They're in the hands of the Technical
      Basis Document team right now going through that
      information. So we're working each one of these
      situations independently and as best we can.
8 DR. ZIEMER:
             Yeah.
9 MR. GRIFFON: Yeah, I just had a question -- or a couple of
      questions.
                  First one's on the telephone interview.
1 MS. ELLISON: Uh-huh.
```

2 MR. GRIFFON: It says 200 to 300 per week. Do you know how

```
many interviewers on average are doing -- are
      conducting the interviews?
3 MS. ELLISON: I might have to ask Dave to answer that.
      know --
5 MR. GRIFFON:
               Ten? Is that the --
6 UNIDENTIFIED: About ten.
7 MR. GRIFFON:
               Ten?
8 MS. ELLISON:
               About ten?
9 MR. GRIFFON: And is there -- has there been any effort --
      maybe Dick can answer this, too, as -- are -- are there
      certain people who are doing certain sites? Are you
      grouping them with interviewers that have expertise or
      knowledge of certain facilities or how -- how is that
      being...
5 DR. ZIEMER: Okay, I think Dick Toohey may be able to
      respond to this. There -- is that mike working? Yes.
7 DR. TOOHEY: (Off microphone) Is this one on?
8 MS. ELLISON:
              No.
9 DR. ZIEMER:
              Is there a mike we can use for --
0 MR. ELLIOTT: He's got it working now.
1 DR. TOOHEY: Okay, yeah, there we go. Okay. Actually we
      have a total of 16 interviewers, but with, you know,
```

vacation, things like that, the average working a day is probably 12 to 14. Our goal has always been to get them to be site-specific. But since we're still in kind of what I'll call a batch mode where we're -we're trying to knock all the Hanfords out, all the Savannah Rivers out, the sites where we have the site profiles completed -- we're not really doing that yet. 8 MR. GRIFFON: Okay. Just one more. On the slide before that, you talked about the data requests to DOE and the percentage of responses, and I think -- I've tried to ask this before and I'm not sure how clear I was, but first of all, does -- do those statistics include data requests for information that might be used for the site profiles, or is that only for personal dosimetry records or ... 6 MR. ELLIOTT: It's only -- those numbers represent the requests that we've sent for personal dose information. The requests that we've made to DOE for site profile is tracked in a separate system. 0 MR. GRIFFON: So anything in here, it's fair to say, is personal dosimetry or medical records or work history records or those sorts of things. Right?

elsewhere.

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1 MR. ELLIOTT: Dosimetry records, not medical records.
      medical. Dosimetry records -- there may be -- in those
      numbers there may be requests for -- if they couldn't
      find personal monitoring information, there might be a
      request for co-worker data or alternate -- you know,
      the hierarchy of data that we seek.
7 DR. ZIEMER:
              Okay. Tony Andrade, and then back to Roy.
8 DR. ANDRADE:
               I just wanted to comment that one thing we
      should not forget is that dosimetry records are
      difficult, in and of themselves, to understand and to
      send back in the appropriate format for dose
      reconstruction, especially in the case of inhalation.
      Way back in the early days we used to have limits
      imposed on us in terms of body burden, and so some of
      the data that were recorded were in terms of
      percentages of body burden. Then in the early 90's --
      well, after World War II and then on into the 90's,
      dosimetry records for, in particular, inhalation
      uptakes were kept in terms of annual doses. So those
      are easy. Those are easy if you're talking about an
      employee at a -- at one facility, who's never worked
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1 When you start to have to add records from places that an
      employee may have worked, then that becomes a little
      bit more difficult.
4 Then after about the early 90's, legislation was passed such
5
      that we had to maintain those type data in terms of
      committed effective dose equivalent. That's a 50-year
             It's basically an exponential function of the
      dose that you will receive within your body of the
      radionuclides as they sit there and decay. So given
      that NIOSH requires these data in terms of annual dose,
      they have to go back and work with that function and go
      back and formulate and formulate it in terms of years.
       So when somebody says oh, how come these people aren't
3
      responding so quickly, just factor that into your
      thinking, especially at Los Alamos where we worked with
      uranium and with plutonium and have kept records for
      people coming in from other facilities. Let me just
      say it's not an easy process to send back dose records
      very quickly for one person.
O DR. ZIEMER: Okay. Comment or response from Larry, and then
      we'll go to Roy.
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2 MR. ELLIOTT: I appreciate Dr. Andrade's comment, but I

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would -- I feel the need to clarify something. We
      don't ask for annual cumulative dose. We ask for the
      raw numbers that were used to build that annual
      cumulative dose, and I think that even goes further to
      add to the difficulty of providing a response to our
      request, because we need -- you know, if the badge
      exchange frequency was a monthly basis, that's what we
      want to see. If the bioassay was done on a quarterly
      basis or an annual basis or what have you, we want to
      see the raw numbers. That's what we're after.
1 DR. ZIEMER:
              Thank you. Okay, Roy.
2 DR. DEHART:
              In the presentation you indicated that the
      final dose reconstruction was sent to the claimants
      approximately -- a little over 1,000. What kinds of
      responses are coming in from those claimants who are
      receiving those doses?
7 MS. ELLISON: In the finals?
8 DR. DEHART: On the finals, yes.
9 MS. ELLISON: As far as I -- once -- it's a final copy of
      the report to let them know that it has been forwarded
      then to -- to the Department of Labor for final
2
      adjudication.
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1 DR. DEHART: Are they making any comments in -- back to you
      as to correct or --
3 MS. ELLISON: Not on finals that I'm aware of. I would
      think minimal -- you may --
5 DR. ZIEMER: Yeah, let me ask either Larry or Dick Toohey --
6 MS. ELLISON: -- need to ask...
7 DR. ZIEMER: -- also to address that. I think they do have
      the opportunity to comment on that, so what -- I think
      that's the question, are we -- are we in fact getting
      comments and what are the nature of those.
1 MR. ELLIOTT: Yeah, and I think Dr. DeHart's question is on
      the OCAS-1 stage --
3 MS. ELLISON: Right.
4 MR. ELLIOTT: -- because that's the stage where we send a
      draft dose reconstruction report to the claimant with
      this OCAS-1 form that they are asked to sign off on,
      which imparts that they have no further information to
      provide and allows us then to move the complete dose
      reconstruction over to the Department of Labor. We do
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hear comments back at the OCAS-1 stage. A lot of

people -- it runs the gamut from thank you, I finally

have an answer, to I don't agree with what you've done,

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to writing things on the OCAS-1 that have no bearing on
      the case but we have to take that into account.
      this is all captured and tracked in the administrative
      record, so it runs the gamut. I would offer that the
      majority of OCAS-1's that are returned to us are simply
      signed off on and -- and that's it. We don't have any
      further -- further commentary that is provided at that
      point.
9 DR. ZIEMER: Okay. Mike Gibson.
0 MR. GIBSON: Just another comment that complicates I think
      the issue of the dose reconstruction is at least some
      of the sites, if not all, when the records were kept
      some years ago it was just recorded in gross alpha, not
      specifically the radioisotopic -- that was -- that was
      used, so it could -- it could make a difference in the
      dose consequence.
7 DR. ZIEMER:
              Thank you. And certainly the internal dose
      issues are complex, depending on the site.
9 Any further comments?
O MR. ELLIOTT:
               Chris has a little more to show.
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1 DR. ZIEMER: Okay, Chris, you have some additional --

2 MS. ELLISON: I have one more piece --

1 DR. ZIEMER: One more piece.

2 MS. ELLISON: -- if you'll bear with me. Currently on our web site we have a claim information page, and if you have looked at it, it gives running totals for the various steps in our dose reconstruction. Our -- the director of NIOSH, Dr. Howard, has requested and wanted to see specifically of the cases we have in-house just where are they in our system. So we've developed a flow chart and we're currently working on developing it and getting it on the web site, and I plan on sending screen shots to the Board once it's a little bit further along, so you'll see this before it goes up on the web site.

4 Basically it breaks down the cases that we have in-house and shows which individual step the cases are in. And with each of the steps, you can click on the boxes and there will be a breakdown then of the information contained on the site. The gather exposure information -- I'm having a hard time getting it to scroll down for me, there we go. Oh, it does break -- we have most of the information broken down into the four district offices that we receive the information from, but we're hoping

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that this will kind of show the public a little bit
      better specifically where that 14,000-plus cases are in
      our system, how many of them are actually waiting on
3
      the exposure information, how many of them are at the
      telephone interview stage. And that's primarily just
      what I wanted to show you, the draft DR's and things.
7 Any questions?
8 DR. ZIEMER:
              Jim?
9 DR. MELIUS:
              Yeah. I don't think most people out there
      really care about the DOL district offices.
      any way you can break it down by site? I know people
      work at more than one site, but how you count that gets
      a little complicated, but it certainly would be a more
      meaningful --
5 MS. ELLISON:
              Okay.
6 DR. MELIUS: -- number for people to --
7 MS. ELLISON: Like I said, right now we're working on that -
              I understand, I'm just saying --
9 DR. MELIUS:
0 MS. ELLISON: -- (Inaudible) right, and those are the type
      of comments we're looking for.
2 DR. MELIUS: 'Cause I think that would be -- help -- help
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people, and if there's a way you could array the -- a screen that would have the -- the different steps and the numbers and just as a table that people could look at quickly by site, maybe it's another -- another page or something that would --

6 MS. ELLISON: Okay.

3

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7 DR. MELIUS: -- with the sites down the left-hand column --

8 MS. ELLISON: Thank you.

9 DR. MELIUS: -- I think that would be helpful.

O DR. ZIEMER: Thank you, good suggestion. Any other comments or suggestions? Henry?

2 DR. ANDERSON: Just one. That would be -- we're on our number 19 meeting, and I guess one thing that would be helpful is it's nice to see the numbers changing and advances being made, but it's hard to come up with an overview as are we catching up, are we further -falling further behind. The process seems to be now kind of in play, the number of requests coming in are -- appear to be going down, but with 504 in the last quarter, it just -- grossly it appears to me as though we may be falling even further behind, so it'd be nice if we could have some kind of a summary of what changes

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have been made, you know, what new is coming in --
2 MS. ELLISON: Uh-huh.
3 DR. ANDERSON: -- and you know, how many we -- we can see
      how many are going out, but where do we stand on the
      overall program of -- can we say we're starting to eat
      into that backlog?
7 MS. ELLISON: Okay. Thank you.
8 DR. ZIEMER: Other comments, suggestions?
9 MS. ELLISON: And as I said, I plan on sending screen shots
      of the various screens for further comment and for
      further review.
2 DR. ZIEMER: Thank you.
3 MS. ELLISON: Uh-huh.
4 DR. ZIEMER: Okay. We appreciate your presentation, Chris.
5 MS. ELLISON: Thank you.
              STATUS REPORT - DEPARTMENT OF LABOR
7 DR. ZIEMER: The next item on our agenda is a status report
      from Department of Labor. Jeff Kotsch is here this
      morning with us and he'll make that presentation.
      Jeff.
1 MR. KOTSCH: I think it's on, isn't it?
2 DR. ZIEMER: Yeah, it's on.
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1 MR. KOTSCH: Good morning. My name's Jeff Kotsch. I'm the
      health physicist with the Department of Labor's
      program. Pete Turcic was unable to attend this morning
      because he has a DOL management meeting this week, but
      we pretty much promise you'll see Peter next time, so
      hopefully that'll be the case.
7 We're -- all of this data is pretty much current as of
      November 27th of this year, and one problem that we
      wrestle with is always trying to synchronize with --
      trying to get our numbers to kind of match NIOSH
      numbers, and I know it drives my bosses, both Pete and
      Shelby, crazy when we can't come up with the exact
      numbers, but I don't know that we'll ever get there,
      but we'll -- I think we're fairly close. But you just
      have to keep that in mind.
6 The total number of claims received to date or as of
      November 27th is 49,113. The first five categories up
      there are primarily the ones that are covered under the
      statute, and the last one, the other, is -- is the ones
      that are not. Those are the respiratory conditions
      that we get -- the COPD's, the asbestosis, the heart
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conditions, things like that -- things that are not

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covered by the statute.
2 The bulk of the ones that are covered are primarily cancers,
      33,766, and then the rest are -- as indicated there --
      the beryllium sensitivities, the CBD -- the chronic
      beryllium disease, the silicosis and the RECA
      compensation that's part of the Department of Justice
      program.
8 Now one thing during this presentation, we're shifting
      between cases and claims, and just so you get -- have
      to keep in mind again that the cases -- there's one
      case for every employee, but there's -- and if the
      employee's still surviving, he's the claimant, but if
      he's deceased then you either have a spouse or one or
      more children as a claimant so the claimant number is
      always -- or the number of claims is always greater
      than the number of cases.
7 So there's just, reading left to right, the case status in
      the Department of Labor, 14,838 cases that were sent to
      NIOSH as of November 27th. Recommended decisions were
      issued by the four district offices, a little over --
      about 21,400. And if you go over -- well, I'll just go
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      left to right. Pending final decisions, about 1,500.
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Those would be in our FAB -- the final adjudication
      branch -- in the four district offices, as well as our
      national office. And then the FABs also issued about
      19,900 decisions.
5 Pending action, the second from the right, about 1,500 cases
      are still pending some kind of review in our district
      offices. And then the total number of cases is 37,192.
8 For the final decisions, the claim approval is 10,729 and
      denial is 14,324, and then the recommended decisions
      are about 11,200 for approval and claims denied of
      about 16,500. Again there's the NIOSH -- the number of
      cases sent to NIOSH, the payments issued as of November
      27th are 9,483. The amount of compensation is $700
3
      million 474,000 or 475,000. The amount of medical
      benefits paid is $21 million about 205,000. And
      there's a summary of the total claims, the total cases.
       The initial decisions, these are the point at which we
      either -- when we consider initial decisions, the point
      where we either have gotten to the point where we send
      the case on to NIOSH for dose reconstruction or, if
      it's not a cancer case or it's something we can
2
      continue adjudication on -- the SEC cases, the
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beryllium cases, the silicosis cases and the -- and the ones that are covered under the (Inaudible), but any one that referring for -- I'm sorry -- so anyway, recommended decisions for 27,000 about 700 claims, recommended decisions for the -- of that 21,000 about 400 cases. So at that point we're about 96 percent completed as far as the process goes to get to the initial decision, either forward it to NIOSH or just to continue with the recommended decision. O Again there's a summary of the claims and the cases. final decisions, with 25,053 claims of which there's 19,835 cases. And there's the percentage of final decisions, about 53 percent. And the bulk of those would be the NIOSH cases that are still awaiting some decision as far as going to a final decision. 6 That's just the breakdown of the way the claims are. the 25,053 total claims distributed (Inaudible) down 10,729 finals approved, 14,324 denied. The bulk of the denials is that purple column there. Those are the non-covered conditions -- the respiratory, the heart, the other types of conditions that are not covered under the statute. And then in decreasing order, the -

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- 2,318 for the employees that are not covered at -- or
      worked -- had worked at covered facilities, ineligible
      survivors, conditions not related to employment,
      insufficient medical evidence, and then cancers not
      related with POC's less than 50 percent.
6 As far as processing time, I think the last time we went
      through by quarter the processing times for the last
      year and we saw that as far as the Department of Labor
      that the percentages were coming below our target goals
      and continue to be that way. For the last quarter the
      average initial processing time for the AWE, beryllium
      vendor, DOE subcontractor claims is 103.5 days versus a
      goal of 180 since we assumed that it's harder to --
      it's -- it has taken longer to get that information.
      The average initial processing time for the DOE and
      RECA claims is about 76 days against a target of 120
      days.
8 Status of the referral of the 14,000 -- what we see as the
      14,838 claims and cases that have gone for NIOSH
      referrals as of the 27th of November, the cases --
      we're showing 100 -- or I'm sorry, 1,146 cases returned
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      from NIOSH. I think the NIOSH number is lower.
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processed?

again this is one of our disconnects I know we're working on with our systems people to look at these cases and to see how we report these things between 3 ourselves and how -- I guess also how we're looking at what constitutes cases. And those are the breakdowns for the completed cases returned to NIOSH -- cases with recommended decisions, we have had 863, the acceptance is 321 of those, 542 denied. Cases with final decisions, 478, of which a little more than half, 254, have been accepted as of November 27th. That's the end of this. 2 The one thing I just wanted to comment on is the last time we spoke about -- or you asked questions about -- or 3 DOL outreach activities. Pete mentioned to me -- when I went back I talked to Shelby and Pete. We certainly are doing outreach. We have developed a plan for that and Pete asked me if he could just present that at the next meeting, that would be fine. So if there are any questions? O DR. ZIEMER: Thank you, Jeff. Do you have any specific numbers on the number of SEC cases that have been

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1 MR. KOTSCH: No. I mean I don't -- not with me. I -- this
      is not -- the case numbers are things that I don't
      normally deal with as part of my job, and I'm just
      trying to think back to the statistics I've seen.
      don't know whether --
6 DR. ZIEMER: It seems to me that might be of interest to us
      to -- maybe next time --
8 MR. KOTSCH: I certainly -- I know we -- I know we break
      those numbers --
O DR. ZIEMER:
             I'm sure you have the numbers. I think it
      would be of interest to this Board to know how many SEC
      cases have been processed and --
3 MR. KOTSCH: Yeah, we can do that.
4 DR. ZIEMER: -- and approved. Leon, a question?
5 MR. OWENS: The initial processing claims time, is that
      inclusive of the time that it takes DOE to do the
      records retrieval, or is that excluded?
8 MR. KOTSCH:
              That includes the time that we get -- well,
      that -- that does include the time for us to get answer
      back from DOE as far -- at least as far as employment
      goes. And certainly -- I mean the NIOSH process is
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      more of -- time-consuming because they're going out for
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2 DR. ZIEMER:

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another document retrieval that's more extensive than
      the one we have. And we also default -- sometimes if
      we don't get a response back by -- from DOE in a
      sufficient amount of time, we'll default to other
      mechanisms like Social Security and things like that to
      confirm employment 'cause we're primarily after
      confirming the medical conditions and confirming
      employment.
9 MR. OWENS: Okay, so in the event that you don't receive the
      records from DOE in what the Department feels is a
      timely manner, then you would resort to other agencies
      in order to try and verify that employment?
3 MR. KOTSCH: Yeah, I mean we continue to ping DOE as far as,
      you know, trying to get a response one way or the other
      from them that they either do not have records or
      they're unable to provide records, and then we'll move
      forward. I mean we continue to seek from them the
      records that -- but we try to get the response back
      from them that they -- that they do not have records
      before we move on to the Social Security or union labor
      -- labor records or things like that.
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Tony and Jim and Gen.

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1 DR. ANDRADE: Once a case has been considered to be at what you call initial decision, what is the time between that point and the point that the case is sent to NIOSH? And also I'm just curious, how is the case sent to NIOSH? Is it just a direct digital transfer or some other means?

7 MR. KOTSCH: No, the -- what -- when a case -- for the NIOSH cases, basically once we've developed the information on employment and medical condition, there's no real -as soon as that's assembled, that information is now transferred to NIOSH. There's no -- I don't -- I don't know how to ascribe a time to that. When it is transferred, it's unfortunately transferred as a hard copy. The case file is copied by Department of Labor and a copy goes to NIOSH. We have looked in the past to -- especially with the NIOSH referral summary document, which is our basically summary of the case, transferring that to NIOSH digitally so at least that would -- that could -- that could be done and we're still looking into that. But our two systems unfortunately are not highly -- our computer systems are not really compatible with one another.

1 DR. ZIEMER: Jim?

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Going back to the SEC case issue, one of the 2 DR. MELIUS: things that we talked about many meetings ago and I think it may be useful to tie together in this context is that there are also a number of cases that -- from the SEC sites that don't meet the criteria in terms of the amount of time worked, that sort of overlap between the SEC program and the NIOSH program, and at some point there's some issues with how -- how those are going to be dealt with that we have decide on -presumably also at some point Ted Katz'll finally get his job done and we'll have SEC regs out there and we'll be able to -- I think that's one of the issues we have to deal with, so it may be a way of tying this all together that I guess is what I'm proposing to -- that we as a Board need to be thinking about, and NIOSH does, also. And I think it would be helpful if we knew the numbers involved in this overlap area and in some of the situations so we can sort of think about how to -- how to approach it.

1 DR. ZIEMER: Jim, you're asking about the numbers who don't meet the time requirement at the SEC sites who may have

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submitted for a claim anyway or --
2 DR. MELIUS: Yeah, they've submitted for a claim, they don't
      meet the time requirements --
4 DR. ZIEMER:
              Time requirements --
5 DR. MELIUS: -- for the SEC sites --
6 DR. ZIEMER: Which is basically the 250-day -- yeah.
7 DR. MELIUS: They fall in between the sort -- the --
8 DR. ZIEMER:
              Yeah, I understand.
              -- you know, are they an SEC, we don't -- we
9 DR. MELIUS:
      can't really deal with this till we have the SEC regs
      out, but at that point sort of having the numbers and
      understanding the numbers involved, the situations
      involved, I think may help in terms of dealing with
      this issue.
5 DR. ZIEMER: Larry, response?
               So you're asking, Jim, for the numbers of
6 MR. ELLIOTT:
      cases that were submitted but didn't qualify under the
      SEC.
9 DR. MELIUS:
              So they come to NIOSH --
0 MR. ELLIOTT: They come to NIOSH for dose reconstruction.
1 DR. MELIUS: Reconstruction, and there's this issue of how
      do you account -- how are we going to do their dose
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reconstruction and...
2 MR. ELLIOTT: Well, we are doing dose reconstruction for
      those cases. We've actually returned several cases for
      Piketon, Paducah and K-25.
5 DR. MELIUS: And we need to look at that issue. That's what
      I'm saying.
7 DR. ZIEMER: Gen Roessler.
8 DR. ROESSLER: Jeff, on your slide number four you gave some
      dollar numbers. It was $700 million paid out in
      compensation and about $21 million in medical benefits.
       I think -- I'm surprised that the medical benefits is
      a small fraction of the total compensation and I'm
      wondering what the reason for that is, a number of
      claimants have died or ...
5 MR. KOTSCH: Well, part of it is that if you're -- if you're
      in a survivor condition, there's no medical payments.
      It's just a $150,000 compensation. If you have an
      employee who's surviving, then he will submit for
      medical benefits. Early on we were surprised that they
      were not submitting for medical benefits.
      some problems I think with the health care providers,
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you know, accepting our -- basically our compensa -- or

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trying to come directly to use, that's the way we wanted to work it. We didn't want it to have to go through the -- you know, the employee to pay them. We wanted to do it directly, so there were some -- some things there that had to be corrected to move forward. But yeah, we're surprised, too, that the number's a little bit lower than we expect. 8 DR. ZIEMER: Henry? 9 DR. ANDERSON: I don't remember the actual numbers on your slide, but it appeared there were quite a number of

cases where the medical records were insufficient or something like that, and I was wondering what -- what -- what other kind of things that are insufficient? individual has said that they had a disease and then the record review couldn't document that, and what's kind of the step after that, is -- you know, what further do you do if -- if they say they have it but the hospital has destroyed records or something?

9 MR. KOTSCH: Yeah, and that's primarily I think what the I know I'm looking at a case right now where there's -- you know, the physician came back and said I don't remember exactly treating your husband back in

1970 and my office destroyed all the records, you know, after they're ten years old kind of thing. Then you're left basically with -- you know, if the doctor doesn't remember -- the physician doesn't remember, and some do write fairly extensive letters that, you know, well, I don't have records, I -- you know, remember that, you know, I worked with this patient and -- and he had these kinds of conditions over this pa-- you know, over those years. All we can do is ask for affidavits or try to get other information from the medical people. Occasionally -- and we're always amazed that sometimes people have kept their records, even though they've been destroyed by the hospitals but they personally have kept them, which has really been of benefit to them, obviously, when the records unfortunately no longer exist, you know, in the hospital. But yeah, it's a difficult thing for us and -- to try to find -or help the claimant find the medical evidence that provides us with something that substantiates the claim.

1 DR. ZIEMER: Jeff, can you speak to us about the numbers of appeals for both NIOSH final decision cases and what I

would -- I'll call right now the non-NIOSH, the ones that don't require dose reconstruction, which would include your -- I don't know, any of the other ones that you're handling, but what's the experience on the appeals -- of the denials? 6 MR. KOTSCH: Yeah --7 DR. ZIEMER: I assume no one's appealing the acceptances. 8 MR. KOTSCH: No, no one appeals the acceptances. I -- and unfortunately, I don't have real good numbers for you. I -- and the only things I really work with, I happen to be the focal point for the technical objections to the NIOSH dose reconstructions, and I've seen about or I have on my desk probably -- I mean have total, since 3 the beginning, maybe 24 or 25 cases that have technical appeals to the NIOSH process. As far as the overall numbers of the objections to the process itself, I'm sure it's much higher because it wouldn't be just -- it would be denials for all the other things and not just the NIOSH cases. We can certainly bring those numbers the next time, as far as both the general objection rate to the FAB decisions as well -- or the recommended

decisions, and as well as the ones that relate to the

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NIOSH (Inaudible) --
2 DR. ZIEMER: Well, I don't know if one distinguishes between
      objections and formal appeals. I mean someone may
      object, but I'm asking --
5 MR. KOTSCH: The appeal -- appeal process is on the final.
      They have an opportunity at the recommended decision
      stage to object, and then it goes forward.
      included in -- by FAB into --
9 DR. ZIEMER:
              Is that, what you're calling an objection, kick
      off an appeal process then?
1 MR. KOTSCH:
             Well, the -- what -- I'm sorry, I should
      clarify myself. The objections -- when I talk
      objections, I'm talking mostly at the recommended
      decision stage. And then they can go into the -- that
      can be factored into the final decision and at that
      point is kind of what the appeal process is -- there's
      a couple of elements --
8 DR. ZIEMER:
             Right.
9 MR. KOTSCH:
              -- you know.
O DR. ZIEMER: And that's what I was asking you about. Maybe
      someone can let us know next time how that's --
2 MR. KOTSCH: Yeah, we can (Inaudible) as far as the fin-- we
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-- all the things I'm looking as far as technical
      objections are at the recommended decision stage.
3 DR. ZIEMER: Roy, you had a question?
4 DR. DEHART: Actually it was just an expansion on the
      medical record or lack thereof that we're seeing.
      panel of three physicians reviews -- currently, at
      least -- the information that's provided from the
      Office of Employee Advocacy, and we have seen records
      that have come to us that have no medical documentation
               There is no diagnosis that's documented.
      There is no record of treatment or management.
      are typically elderly people who have passed away and
      the relative is filing on their behalf, and they have
3
      no access -- they don't even know perhaps what hospital
      or what doctor. And of course there's no way we can
      move forward with anything on those kinds of -- that
      lack of information.
              Those are for the subpart (d) cases --
8 MR. KOTSCH:
9 DR. DEHART:
              Yes.
O MR. KOTSCH:
              -- but yeah, admittedly we see the same thing
      in the subpart (b) cases.
2 DR. ZIEMER: Further questions? Thank you, Jeff.
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1 MR. KOTSCH:
              Okay.
2 DR. ZIEMER: Appreciate your input this morning.
3 We're a little bit ahead of schedule, but I think we'll go
      ahead and take our break. Let's -- we can take a
      little longer than the 15 minutes, maybe about 20
      minutes or so, then we'll reconvene. Thank you.
7 (Whereupon, a recess was taken.)
               SITE PROFILE STATUS AND ROLL-OUT
9 DR. ZIEMER: We can proceed. The next item on our agenda is
      site profile status, and the -- I'm going to sort of
      say pinch-hitter, but he's very well qualified, Stu
      Hinnefeld, who works with Jim Neton on these activities
      from NIOSH -- and Stu, we're glad to have you here
      today, and Stu will present the site profile status and
      roll-out.
6 MR. HINNEFELD: (Off microphone) Thank you.
                            (Pause)
                 So that's it? Okay, sorry. I am really
8 MR. HINNEFELD:
      substituting for Jim Neton today. He was unable to
      travel this week and so I'm here in his stead, and I'm
      here to present sort of an update on the presentation
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that he presented at the last meeting about site

profile status and the progress that we're making on preparing the site profiles for quite a number of sites, actually. 4 Sort of recapping the information that Jim presented in St. 5 Louis, and also presenting for anyone who hasn't previously seen this or heard about this, site profiles are documents that are used by dose reconstructors to provide consistent interpretation of the information provided from the various DOE sites so we have a consistent understanding of what the bioassay data means, what their bioassay or what their external monitoring technology was, and that allows us to provide consistent dose reconstructions for a particular site. Each particular profile will address one site and one -- perhaps one particular type of exposure -- well, actually each document will. Several documents will be rolled into one site profile. Ιt helps us to minimize individualized interpretation. In other words, we can have a consistent set of rules for interpreting the data. And it's used as a handbook by the dose reconstructors to provide -- to guide them in 2 their work. And they're intended to be dynamic.

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learn more about various sites and their approaches and
      their technologies, then we may in fact modify the
      information in the profiles.
4 We are publishing our completed profiles, including the
5
      individual pieces of profiles, which we call Technical
      Basis Documents, as they're approved, on our web page.
       There is a little bit of a administrative process that
      happens after one is initially approved.
      little time lag between the approval and appearing on
      the web page, but they're all being placed there.
      We're encouraging comments on these, and if anyone
      feels compelled to comment on the -- either the content
      or the proposed application of these site profiles,
3
      those comments could be submitted to the NIOSH docket,
      and there is a specific docket established for each of
      the approved documents. So again, if you go look at
      them on the web site, it's fairly apparent there's a
      link to -- to what has been -- what has been provided
      to the docket for that -- for that particular document.
0 We are arranging to present the completed documents as
      they're being completed to union representatives and
      other interested parties in the vicinity of the
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affected site. One of those briefings has actually
      been done since the St. Louis meeting. It was done in
      November at the Savannah River Site, and the next one
      is scheduled for Hanford in January.
5 And there's information about the team members for the teams
      of consultants who compiled these initial versions of
      the profiles on our ORAU -- our contractor's web site.
8 Now in order to contact the docket or submit comments, here
      are the ways that you can contact the NIOSH docket
      office. Written hard copy comments can be sent by mail
      to the address here. We have telephone and FAX numbers
      and e-mail that then provides comment to our docket
      office, and we'll know it is a comment to the docket by
      using that e-mail address.
5 I might mention that when I was looking at my printed -- the
      printed copies of my slides, there are a number of
      spaces that found their way into the slides that don't
      appear on the slides themselves. And not being very
      good at this particular software, I apparently wasn't
      able to get them all out of there, so I apologize for
      the printed copies of the handout.
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2 Our latest status on the site profile status, this again is

essentially unchanged from the St. Louis meeting. There are 15 facilities we are working on at the same time. We expect to complete them by the end of the calendar year, and that's getting close. There are very, very many documents that are -- have been reviewed and are in comment resolution and are very close to being approved. And I did not go back and update the percentage of claims, but it's our expectation that these documents will cover very close to 77 percent, or something close to that number, of the total number of claims. In other words, the total number of claims we've received we have to do dose reconstructions from, some 77 percent of them came from this first 15 or so DOE facilities we are now working on. 6 And in addition, since the St. Louis meeting or the last Board meeting, we have completed a couple of complexwide approaches for processing dose reconstructions. One is for DOE facilities and one is for AWE facilities. These are somewhat limited in their applicability. They obviously can't be applied to all 2 DOE claims or all AWE claims, but there is a set of --

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a set of claims that do lend themselves to a complex-
      wide approach for dose reconstruction, and I'll speak
      more about that in a little while.
4 Site profile status, this was as of November 24th, which was
5
      when the -- I had to complete the preparation for the
      presentation. You can see two of the larger DOE sites,
      Savannah River Site and Hanford site, those site
      profiles are complete. And then for several other
      sites there are either one or two pieces of what I call
      here as a five-piece site profile. There are about
      five Technical Basis Documents in each site profile,
      and then there is a sixth section called an
      introduction, so I don't really count the sixth as one
      of the research-oriented pieces of the document, so I
      entered these as five.
6 After this slide was prepared, a third Technical Basis
      Document from Y-12 plant was approved, as well, so
      that's the change as of late last week from this slide.
       And you also note down here we've completed a DOE
      complex-wide approach.
1 For AWE sites, I believe this is the same list that was
      available at the last meeting, except for the addition
```

of the complex-wide uranium AWE facilities. 2 Since we're in Nevada, in the vicinity of the Nevada Test Site, I researched where we were on the sections of the Nevada Test Site profile, the various Technical Basis Documents that comprise that document. The profile initially will consist -- here I go back to the six sections, that sixth section being the introduction, and then the other five sections are site description, what we call occupational medical exposure or X-rays received as a condition of employment like during an annual physical, internal exposures, environmental exposures -- those are all fairly far along and in review and comment resolution. The external dosimetry section is yet to get into the comment resolution stage. It's in our contractor's review stage. And then of course the introductory section, which is sort of just a summation of the information of the other five. 9 Okay, the -- I want to spend a little time describing the 0 complex-wide DOE technical basis. We also did a complex-wide AWE technical basis for a limited set of 2 atomics weapons employers that met certain conditions.

First of all, they had to -- their AWE work had to be only with uranium. If there were any other radionuclides associated with their AWE work, then they 3 would -- that -- that particular site could not be -claims could not be processed from that -- from that site through the complex-wide approach. They typically would expect to have a fairly limited scope of AWE work, not a site that did -- cranked out tons and tons of uranium years after years after year. And so it -the complex-wide AWE approach takes some very conservative, in our terminology -- in other words, very high potential exposures and essentially says even under these conditions for this certain set of cancers, it looks like these -- these claims won't be compensable even under these worst-case conditions and therefore it allows some processing of some AWE sites claims. 8 For the complex-wide -- the complex-wide process is what we refer to as an efficiency process which follows from the regulations statement that dose reconstructions done under worst-case assumptions, and if you do a dose

reconstruction under worst-case assumptions and it's

clear that the probability of causation would exceed 50 percent, then in those cases you have -- those cases can be considered complete. No additional research will change the probability of causation determination -- or if additional research would only cause the probability of causation to go lower -- and therefore there is no need to pursue and research in greater depth this particular dose reconstruction. that in mind, there are -- there is a population of claims that it would appear would fall into the category where certain worst-case assumptions can be made. And even if those worst-case assumptions turned out to be true, which in many cases they seem almost incredible, that even if they were true that this -this -- the probability of causation on this case -- on this claim will not rise to the 50 percent level and therefore these worst-case assumptions can be used in application to these various claims in order to complete the dose reconstructions and provide answers to claimants who have been waiting quite some time for their answer to their compensation claim.

2 So that was a brief discussion of the purpose of the

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complex-wide AWE -- or complex-wide DOE technical basis
      approach. And it's structured in four documents that
      are called Technical Information Bulletins.
      address the four major types of exposure that you would
      find in a site profile. The ones that would not be
      described are facility and processes, which is one of
      the five -- five topics in a full site profile, and
      then the introductory section, which is the sixth
      section of a site profile.
0 So these are the documents that comprise it. These are the
      first two about internal dose estimates; external dose
      estimates -- you'll notice this is for
      thermoluminescent dosimeters; occupationally-related
      diagnostic X-rays; and occupational doses from elevated
      ambient levels of external radiation, which we
      oftentimes call environmental or occupational
      environmental dose.
8 So describing briefly the approaches that are followed on
      this -- in this particular regimen for dose
      reconstruction, first there is a case selection
      criteria that you limit the applicability of this,
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first of all, to more recent employment. It wouldn't

be appropriate to use a complex-wide overestimates

because we've made certain assumptions about -- in our

process here that would apply to more recent times, say

from 1970 or 1980 forward, but would not necessarily

apply to very early work. So the applicability of the

complex-wide regimen is really limited to more recent

employment.

doses in order to provide confidence that we really
have captured the worst case that this person may have
been exposed to. Use a maximum credible undetected
intake or a implausible undiscovered intake, depending
upon the terminology you want to use, to evaluate a
worst-case assumption for an internal dose. And then
we choose parameters that maximize POC both by
maximizing the dose and by the selection of the
radiation types and photon energy types.

(Pause)

obtay, so for a little more information about the approaches
that are taken here in terms of the maximizing doses
and maximizing probability of causations, I made a few
notes to go through the various approaches to kind of

describe what was done in the various Technical Information Bulletins to describe these -- this maximizing approach. 4 For the internal dose assessment component, first of all, 5 the employment has to be from a DOE site or a national laboratory that had an established radiation control program. It certainly wouldn't be appropriate to do this with -- this approach with an AWE site which had a much more limited radiation protection program because there's certain assumptions about what could or couldn't be seen at the time of employment. hire date for all these claims must be 1970 or later, and for some applications at some sites, 1970 is also a little early and so those cases can only start if the employment -- dose can only be done under this regimen if the employment started later than that. 7 For internal dose assessment, the claims must involve a cancer of an organ or a tissue that does not concentrate the radioactive materials that the person might have been exposed to. The cancer causation or the dose to an organ from internal exposure depends 2 quite a lot on whether that organ concentrates the

radioactive material or not, and if it doesn't, the internal doses tend to be relatively minor compared to organs where it does -- where the material does concentrate. So this approach can only be used for those organs that don't concentrate the radioactive material. 7 The claims should involve people who either weren't monitored for internal exposure or who were monitored for internal exposure and had no positive bioassay result. And they should be for people whose jobs appear to be -- have an unlikely potential for significant internal exposure. So we're selecting a certain population of claims where we're liable to have low external -- low internal exposures -- internal exposures. 6 Okay, the maximum or the -- the implausible undiscovered intake is based on the control concept of maximum permissible body burden, which Mr. Andrade referred to earlier, because that's the way the standards were written in the 70's and 80's, and it essentially -- the worst case assumption for the internal exposure is that 2 the energy employee was exposed to a -- an intake that

would cause a significant fraction of the maximum permissible body burden for an entire list of radionuclides on his first day of employment during his first year of employment, and that based upon the selection of the case, that this had to be from a case with a radiation protection program. Our belief is that the radiation protection program would not miss that sort of an intake, so the maximum credible intake is developed in that way. It is -- the most soluble form of the radionuclide is chosen for the dose reconstruction because that will provide the highest dose for these non-metabolic or non-concentrating organs. And the actual intake is several-fold times the maximum permissible body burden that was in use at the time because of the rapid clearance from the shortlived components -- or short-lived compartments in the model.

8 There are actually two lists of radionuclides that are used in this postulated intake. One is for sites with reactors and one is for sites without reactors. addition, for uranium sites the ten percent maximum permissible body burden provided really fairly low

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chronic exposure could -- over an employment period could achieve ten percent of maximum permissible body burden, which interestingly enough doesn't seem to be the case in very many of the other radionuclides, so the uranium number was sort of artificially inflated, so the postulated uranium intake is quite a lot larger than -- than it would be based on the original calculation method in order to demonstrate that this would be quite a large chronic exposure over a long period of time, and it would still have to have been missed by this radiation protection program. are the key features of the internal dose assessment approach.

4 The elevated ambient level of external radiation approach or the environmental occupational dose approach is based on a review of environmental monitoring reports, as well as some of the site-specific research that's going into the preparation of the site-specific Technical Basis Documents. And what we can find from the review of the environmental monitoring reports is from 1980 on, environmental releases were just not really all that big that they would be causing measured --

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measurable radiation exposures to a radiation monitoring device. From 1970 and 1980 in certain cases at certain sites, from our research, it appears to be the same, as well. And another aspect of this -- this particular component of dose is it's appropriate to add this dose to a dose reconstruction only when this dose is not measured by the person's personal monitoring So the person is wearing his dosimetry device and is exposed to this ambient elevated radiation level while he's at work, his dosimetry device will record The only reason -- and so it would be in that dose. his dose record, unless there was a control dosimeter that was used for a background subtraction that was located in a similar area. So if the control dosimeter was irradiated to this elevated level, as well, and if in fact the site was using that control dosimeter as a background subtraction on their personnel dosimeter, then that excess ambient dose would be subtracted off. And so from our research we found certainly at some of the bigger sites where we're further along in our research that the releases were getting quite small in the 70's and the dosimetry practices were such that

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those -- that inappropriate control subtraction --
      background subtraction was in place. And so we should
      be able to identify additional time frames in the 70's
      to use these -- we think for most of the 70's,
      probably.
6 The medical occupational exposure was established based upon
      an evaluation of literature searches of exposure
      techniques and resulting exposures. Over history
      various studies were done at various times.
      numbers -- there's a particular table of numbers that
      are used for organ doses like -- that would likely
      result from exposures before 1970, another table for
      1970 to 1985 and then another table from 1985 forward.
       These generally reflect the improvement in the
      understanding of the role of filtration, columnation*
      and technique factors and the general improvement that
      can be seen in these various scientific reports that
      were written over time to describe medical dose re-- or
      the medical exposures.
O Since the remainder of these types of radiation exposure can
      really only go from 1970 and forward, the pre-1970
      number in this document won't be used in this complex-
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wide, but they may be used to facilitate the preparation of other medical profile information later on when we can't find specific information on a given site going back before 1970. 5 Finally, for the external exposures, we again have case selection of cases where the radiation exposure appears to be relatively low and the cancer is in a location that is not -- does not have a particularly high risk factor associated with radiation exposure. So we go in by selecting cases that look as if they have a low chance for exceeding the 50 percent probability threshold, select those cases and the external dosimetry must be done by thermoluminescent dosimetry rather than film. And the person's -- should have no neutron exposures. He should either be unmonitored for neutron exposure or he should not have a measured neutron exposure, and he should have a job that would make it look like he probably wasn't exposed to neutrons except maybe incidentally on occasion, but no appreciable neutron exposure. 1 So in this approach we apply an overestimating conversion 2 factor to -- and the purpose of that is to provide an

upper bound for the uncertainties associated with some of the things that were going on in dosimetry technology. For instance, dosimeters respond to different kinds of radiations and different kinds of energies in different fashions. Calibration methods sometimes varied from (Inaudible) calibrations to (Inaudible) calibrations. Workplace radiation fields could be mixed, fields -- not nice clean AP fields like you see in the work -- in calibration facility, and various facilities might have different administrative practices for when you record a dose and when you write down a zero and things like that. 3 So the overestimating correction factor is developed from really two -- two major components. One is a combined uncertainty associated with geometry and calibration, uncertainties associated with measurement at the time. And an upper bound on the uncertainty that was probably being experienced by these sites that were monitoring with TLDs in the 1970's and early 80's. And that's a pretty -- that's established in the Technical Information Bulletin as being about a number of 1.8,

and then there's a maximum organ dose correction factor

to convert this -- this dose number, the recorded dose number as adjusted to the dose to the organ in question, and that's just universally chosen with this -- with this approach to be a maximizing 1.1, which is actually higher than any of the DCS for this type of radiation and this -- this dose conversion factor. those two maximizing values, when they're combined together, give a -- conveniently give a number of about two, so the consistent or overestimating correction factor is a factor of two applied to the recorded dose from these various sites for these selected cases in order to compensate for the uncertainties that may be there, provide an upper bound for what their exposure may truly have been. And then from missed dose standpoint, since the person quite likely wore some dosimeters that read zero, the missed dose concept or missed dose approach on this maximizing approach is just to generate a missed dose as if you wore 12 dosimeters in a year and the limit of detection was 30 and they were all zeroes, and then the -- and apply that dose correction factor again, as well, so double that number as well to arrive at a -- a missed dose for

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these approaches. And the -- this is then applied in a
      -- in the lognormal distribution technique that's
      described in our implementation guide, the limit of
      detection divided by two times N as the geometric mean
      of a lognormal standard deviation.
6 So those were some of the more -- some of the more graphic
      details or bloody details of the approach that was
      using -- that we're using on this complex-wide
      approach. Again, this is for a -- for a limited set of
      claims, and it's to facilitate our ability to provide
      more timely answers to claimants who have filed a claim
      and they deserve an answer to their claim.
3 We also recognize that much of the profile work so far has
      been done from a particular point of view, not so much
      from the affected employee point of view, so the -- at
      the -- this was pointed out at the St. Louis meeting,
      and so we are engaged in processes to identify
      populations of workers, whether they be labor --
      represented by labor unions or whether they be other
      affected workers, to provide input to us in the
      preparation of Technical Basis Documents or for
2
      documents that are nearing completion; after completion
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of the Technical Basis Documents, to provide any
      comments on what was prepared to see if we need to
      provide additional information, modify what was placed
      there.
5 I mentioned earlier that there was a meeting held in
      November at Savannah River. We also have one scheduled
      for Hanford in January. We've established a docket on
      our web page for each of the Technical Basis Documents
      so that any comments made on that Technical Basis
      Document will be viewable there just as easily as the
      Technical Basis Document is viewable. And then we are
      looking into other information-gathering approaches.
3 We have an obligation to provide a plan for providing worker
      input into Technical Basis Documents, following the
      Board's recommendation at the last meeting, and while
      we've not finalized that plan, some of the components
      of the plan will follow along these bullets that I have
      here on the -- on the screen.
9 I'll be glad to entertain any questions or comments.
O DR. ZIEMER:
              Thank you, Stu. Let me ask if you or one of
      the staff can tell us what the response was to the
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November 11th meeting at Savannah River in terms of

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input from people on the site.
2 MR. HINNEFELD: Well, I only know what Jim -- I only know
      what Jim told me. Jim has heard a couple comments that
3
      it was very well-done and thanks for coming and gee, we
      really are glad to hear that. And then there have been
      other comments made in other avenue -- agendas that
      really wasn't what we needed, that wasn't what was
      intended. So I don't know whether different attendees
      came away with different views or how that came about,
      but Jim certainly did at the -- at -- my understanding
      is at the meeting itself, at the end of the meeting,
      the participants who spoke were appreciative and
      thought that it had been -- been done pretty well.
4 DR. ZIEMER: You have some idea of what the level of turnout
      was for that meeting?
6 MR. HINNEFELD: Well, that was not a public meeting. It was
      a meeting with labor --
8 DR. ZIEMER:
              Oh --
9 MR. HINNEFELD: -- certain labor officials --
O DR. ZIEMER: -- okay, gotcha. Gotcha.
1 MR. HINNEFELD: -- and there were eight or ten, I think,
      something like that.
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1 DR. ZIEMER:
              Individuals that had been identified as contact
      points --
3 MR. HINNEFELD:
                 Yes, yes.
4 DR. ZIEMER:
              Thank you.
5 MR. HINNEFELD:
                 Yes.
6 DR. ZIEMER:
              Jim?
              Yeah. I think actually someone -- one of the
7 DR. MELIUS:
      labor officials that was at that meeting will be
      speaking in the public comment period, so we may get
      some additional feedback on that -- on that meeting --
      time.
2 I would like to thank Larry and staff for going forward with
      a plan, and I understand that it's being worked out
3
      still. But appreciate making the effort and setting up
      these meetings 'cause I think they will be -- be
      helpful and I would hope they'd also include some way
      which is admittedly more difficult to deal with some of
      these -- particularly the AWE sites where sort of the
      workplace is closed or dispersed and how do you reach
      out to -- to people, but some sort of a briefing for
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      claimants or something I think might be -- be helpful
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      for -- so people understand what's going on with this
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process.
2 I've got a number of questions, but -- and some of these may
      be more appropriately dealt with later on in our
3
      meeting, but I think the entire Advisory Board did
      receive an e-mail from a person regarding a conflict of
      interest issue on these site profiles at the Rocky
      Flats site profile, and I don't know -- you want to
      take that up later or what the plan --
9 DR. ZIEMER:
              I received an e-mail myself a day or two ago.
      I don't always check my e-mail every day, but I think I
      got it Friday. I think it came from -- perhaps from
      Terry Berry --
3 DR. MELIUS: Yeah, last Thursday is when I -- date off of
      mine.
5 DR. ZIEMER: -- relating or raising an issue that I think is
      what you're referring to. And perhaps -- I don't know
      if this is the time to look at that, but perhaps that
      can be addressed at -- that as a starting point by
      staff. Have you -- you've seen either the -- perhaps
      the staff has not seen the e-mail, but has had some I
      think contact on that issue, have you not?
2 MR. ELLIOTT: Yes.
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1 DR. ZIEMER: And you want to address that now?
2 MR. ELLIOTT:
              Yes.
3 DR. ZIEMER:
              And maybe explain for everyone what the issue
      is and how it's been addressed.
5 MR. ELLIOTT: Yes, Ms. Berry wrote me an e-mail before she
      sent the one to the Board. I have not seen the one she
6
      delivered to the Board yet. But the issue essentially
      is is that an individual on the ORAU team who is
      working on the site profile for Rocky Flats, prior to
      the genesis of this whole program, evidently provided
      some testimony in a litigation on her husband's claim
      and so we're aware now of this. It's actually -- we
      became aware of it once we had the disclosure up on the
3
      web site that this particular individual from ORAU had
      performed in this regard. So we at the Department are
      now looking into this and evaluating what needs to be
      done in this regard.
8 We do take this very seriously and we had another instance
      in -- last meeting in Cincinnati -- or in St. Louis
      where we had a -- another situation called to our
      attention which was slightly different in that a claim
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undergoing appeal process or in the courts, at least,

was -- there was some testimony being provided against the claimant from a principal of one of the firms that our ORAU team subcontracts with, and that particular individual was not serving on any site profile development or dose reconstruction. But we have worked with the ORAU team and that particular subcontractor is being -- will be released. They will no longer be working on our site profiles or dose reconstruction processes. 0 So that addresses both of those that have been brought to our attention. We're still working on this latest one and how we're going to deal with that. 3 DR. ZIEMER: Thank you very much. Jim, did you have a follow-up? 5 DR. MELIUS: I have a number of questions, but just as a follow-up to that particular point, I would just hope as you're dealing with these issues that you're also trying to evaluate other sub -- subcontractors, I guess they would be called, that might have similar problems so that there's some sort of a policy or something being developed so we don't have to sort of constantly

deal with the individual situations. I know it's

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difficult and hard to -- to know when you're somewhat, you know, dependent on what information is provided to you or provided to ORAU and then on to you, but I mean some sort of over -- communication of other subcontractors or something to just make sure that this -- try to avoid this as much as possible I think would be helpful.

8 MR. ELLIOTT: Absolutely, that is part of the review that's underway right now and discussions that are being held and what type of contract language do we need to have in place.

2 DR. MELIUS: Yeah. Yeah. My next question is also sort of procedural, but on Friday I received a FAX of a letter to you, Paul, from Congressman -- Congressman Quinn, Congresswoman Slaughter and Congressman Reynolds from western New York regarding -- or asking the Board to do -- review the site profile for the Bethelehem Steel site and for raising a number of particular questions to -- to address. And this may be more appropriate for us to take up tomorrow, but I just didn't know if everybody else was aware of it on the Board or if we'd received this or --

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1 DR. ZIEMER: I don't know the answer to that. I just myself
      got a copy of that before I left for the meeting here,
      actually studied it on the -- on the plane and I need
      to discuss that I think also with the Department and
      review the related issues to how that particular letter
      might be handled. But I have no knowledge of whether
      other Board members received copies of that letter.
8 DR. MELIUS: Okay. Could we make copies then for everybody
      on the Board?
O DR. ZIEMER: Well, I'll make --
1 DR. MELIUS:
              Yeah.
2 DR. ZIEMER: -- a copy available to Cori and make sure the
      Board has copies of that, that --
4 DR. MELIUS: I've got that with me, so --
5 DR. ZIEMER: Thank you.
6 DR. MELIUS: -- that's fine. I just didn't know what
      happened with that.
8 The issue on these site profiles that we were just briefed
      on, I'm a little -- still a little bit puzzled, I
      guess, as -- or unsure of exactly what the process is
      now. Will we -- we go back a few years when we first
2
      started the advisory committee and we're doing the
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original set of regulations and briefings and so forth -- presented sort of where NIOSH was at that point in time with this program of doing dose reconstructions. We had a number of sort of technical documents that were being developed. And then we -- you moved along and really got the program going, now we've gone -we've sort of changed that original approach. We have the site profile process which originally was -- at least as I remember it, was going to be at the end of the process. You'd compile that from individual dose reconstructions. And now that -- now we're doing it up front and using it as a -- you've described it as a handbook for the people doing the dose reconstructions. And it would certainly be helpful for me and maybe other Board members -- I'd be curious how others feel -- to understand a little bit how your -- how you're using these as a handbook, maybe taking Savannah River or one of the other completed ones and as you've gone through a number of individual dose reconstructions using that, providing with -- us with a briefing at the next meeting on how you are, you know, using that with some examples and so forth. I think --

1 MR. ELLIOTT: Okay. I think the last time Jim talked about this, 2 DR. MELIUS: you really hadn't completed enough to do these -particularly for the more complicated sites, like Savannah River. I think for Bethlehem Steel and some of those, it's more -- more straightforward, but for Savannah River and the other sites, it's a -- more complicated and it certainly would help me to understand what you're doing and for us as part of our review of the program to see how you're doing that, I think it would be useful to do. 2 And secondarily, as part of that, as I recall, when we originally talked about dose reconstructions and yeah, 3 you were going to be developing policies over time, I guess I would see some of these -- both the site profiles, but also these DOE-wide -- industry-wide documents, your guidance doc-- you're developing as being sort of Technical Basis Documents that are part of this process that we'd expect to be developed as you go along. And when we originally started, we talked about the Board reviewing these or sort of having a 2 review process. And I think we as a Board sort of have

to decide -- and you at NIOSH, have to decide how we're going to do this. Some of the early documents got peer review, outside technical peer review. Are we now relying on our -- our contractor to do the technical review on these or is the Board supposed to be doing the review of these, approving these? You know, sort of what is the process going to be? I think, you know, your briefing today is helpful and I appreciate it, but it's not really us reviewing these and --

O DR. ZIEMER: Right.

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1 **DR. MELIUS:** -- in detail and don't pretend to, and I think we sort of have to come to grips with how that's going to go forward 'cause we have a lot of -- lot of your work that figure -- lot -- what you're doing that we just sort of -- we're blinded to. We -- we haven't had time -- and understandably. I mean you've been, you know, trying to get things done, so I don't think it's anybody's sort of fault or placing blame, but I do think that we as a Board have to sort of look at how we're reviewing -- and particularly as we come into doing individual dose reconstruction reviews, we don't want to be in the position of, at a later point in

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time, saying well, gee, this -- this particular overall technical document was wrong or led to serious problems. Now I'm not saying that's going to happen, but I do think we have to talk about that and come up with some way of -- systematic way of approaching this and some of that's I think a better understanding from you. And maybe not at this meeting, but maybe at this next meeting of sort of what -- what are the documents you see being developed, what's changed, what's -- what kind of documents are -- are sort of just procedural, what requires sort of a technical review, then how do we get that technical review done?

3 DR. ZIEMER: Let me ask for other Board members to also maybe weigh in on those comments. While you're thinking about your responses, let me also point out that one of the things that's included in -- at least at the front end of our audit process is to ask our contractor, as they do the various types of audits of dose reconstructions, is in a sense a kind of audit of the usefulness of the site profiles insofar as those assist in the dose reconstruction. So we do -- or I think we're looking toward having in place something

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that will help us do a kind of evaluation because I
      think the process, if it works properly, should point
      out to us strengths or weaknesses on site profiles,
      either generically or individually, as the case may be.
5 I don't recall us -- well, in terms of our charter, we're
      not required a priori to approve site profiles. On the
6
      other hand, the Board itself may decide that it wants
      to look at them in some fashion in the audit process as
      site profiles, or address particular ones. But I don't
      believe our charter calls for us, in advance, to
      approve site profiles.
2 DR. MELIUS: Yeah, I think you're right, Paul.
      think -- well, our charter does calls for review
      individual dose reconstructions --
5 DR. ZIEMER:
              Right.
6 DR. MELIUS: -- to the extent --
7 DR. ZIEMER:
             Right.
             -- they're used there. We also, I think, when
8 DR. MELIUS:
      we approved the original set of regulations that guided
      the dose reconstruction process, we talked about the
      Board advising NIOSH on technical issues --
2 DR. ZIEMER:
              Right.
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1 DR. MELIUS: -- that would develop over time.
2 DR. ZIEMER: Right.
3 DR. MELIUS: Now some of these were technical changes to
      what was originally approved, some of these would be
      sort of further developments. And I just think we have
      to sort of systematize in some way -- part of my
      question comes from sort of what is the full scope
      going to be of our dose reconstruction review contract
      that's out there. Is it going to review every
      procedure? Is it -- you know, do we select? Is it
      every so -- site profile or -- or not, and I -- and
      then we get Congressional letters asking --
3 DR. ZIEMER: Well --
4 DR. MELIUS: -- about that, too, and that makes --
5 DR. ZIEMER: -- keep in mind, too, that an audit process --
      I have to keep stressing this, to myself and to the
      Board and others -- that an audit process is not 100
      percent review of everything that's done.
                                                 In fact, our
      audit process calls for us to review something like two
      and a half percent of the dose reconstructions.
1 Now it's very true that many of those dose reconstructions
      will have used the same site profile for at least part
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imagine that in some form or another we won't in fact be able to evaluate those. But I'm not -- I'm not speaking against systemizing this in some additional way, but simply reminding us that we will in fact have opportunity to address that. 7 I do appreciate learning about what you might call the sitewise (sic) things because there are certain issues that 8 lend themselves to that kind of analysis. those things, whether they're sort of site-wide or more localized, I think will always be subject to interpretation of validity of assumptions. Let's take, for example, the medical exposure of workers. You will make assumptions I think based on practice as to what film speeds are used, what beam filtrations are used, what columnation is used -- all of which affect patient dose to the organ being examined, as well as dose to other organs from either scatter or a practice which in early days was very common and that was to remove the columnation because the lights were not aligned with the true X-ray beam and the way you get -- solve that 2 easily is rather than get the X-ray machine fixed, you

of the process of reconstructing dose, so it's hard to

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just pull out the columnator*. And if your beam is
      wide enough, you'll sure hit the organ you're
      interested in, and every other organ, as well.
4 And I don't know that -- I quess, you know, what assumptions
      are made? Do you make the worst -- a worst-case
      assumption in that situation is that there's no
      columnation.
8 MR. HINNEFELD: As a matter of fact, I think the early --
      very earliest numbers -- table numbers in the medical
      do make that assumption.
1 DR. ZIEMER:
              Okay. 'Cause it was a very common practice.
      Okay. But those are the kind of issues that I -- I
      think a sampling can help us have a level of confidence
      that -- you know, we may not have to sample every
      assumption made, but if you start sampling and it looks
      like the right thing's being done, then that gives you
      a level of confidence.
8 MR. HINNEFELD:
                 Okay.
9 DR. ZIEMER:
              Other -- I don't mean to monopolize this.
      Tony?
1 DR. ANDRADE: Yes, I'd like to just comment that, first of
      all, we should also remind ourselves that we are not --
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we're not an expert board. We are an advisory board, so I'm not sure to what extent we should really look at all of the technical details of even any one of these site profiles. 5 Two is that I do recall that several meetings ago it was announced that site profiles would be developed and 6 that they would be used in a very limited sense. as was presented today, you got a feeling for the limitations that are imposed on their applicability. For example, the age of the employee, the time during which they were working, some of the assumptions made. And I can just imagine the type of filtration or efficiency that you're gaining from these for the types 3 of employees that probably would never achieve a POC of 50 percent. And we're talking about people that perhaps handled the bioassay samples and took them back and forth to the laboratory, administrative assistants that worked nearby to say neutron-generating operations, those sorts of employees in which doses themselves were probably extremely, extremely low. given all those factors -- oh, and along with the fact 2 that we do have a task order out that is supposed to

direct a subcontractor to us to look at -- especially site profiles on top of individual dose reconstruction, I really feel that any further let's say work added to 3 the Board's schedule would at this point just be relatively non-value added and that, insofar as this Board member is concerned, I am quite satisfied with the monthly or however often we meet updates like the one that was just presented that gives me a feeling for how these are being used, what sort of details are going into them and the types of analyses and assumptions being made within them. That's all. 2 DR. ZIEMER: Okay. Let's -- Mark and then Jim. 3 MR. GRIFFON: Yeah, I just -- I do agree with one part of what -- when Tony said that we do have a task order out, and the contractor's going to review the site profiles, even though we're not -- and just a reminder that the contractor's work is -- is the Board's work. I mean the contractor's working for the Board, so we are going to be reviewing site profiles, and I think that's where we're going to get into the meat of it. 1 I do have a couple of comments about the presentation, 2 though, just -- some of which I've probably said

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before. But I -- you know I still -- in your second slide, I see things like limited scope, and I do understand, you know, from your presentation what you meant by that. But the fear I still have with some of what I've seen so far is -- is the idea of -- there are some sites where I think a better understanding of operational details is going to give a very different picture of potential worst-case doses. And if we just skim the surface with a limited-scope site profile -and I'm not saying -- I mean I know it's a lot of work to do these things, too, but we could easily miss, you know, some very -- some operations which have very different exposures than the general building, for instance, and I've found that in some of the work that I've done. And it may not affect a lot of the workers on the site, but it may turn out to be several of the claimants. So you know, without going to that level of detail, I fear that we may miss some of that and underestimate worst-case doses for a certain fraction of people. That's one thing. 1 The other question I -- this is more of a -- go as a statement. The other question I had was later in the

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presentation you talked about the missed dose with the external exposures. I didn't really hear you address unmonitored dose, and I know that's come up again and again during public comments, we've heard it over the years from DOE hearings and things like that. employees report anecdotal reports that their badges were tampered with, they weren't badged for certain high level operations. How are -- how are you handling potential unmonitored exposures or -- in your -- in your...

I guess on the face of it, without looking 1 MR. HINNEFELD: at a specific instance of a specific case and a specific set of claims, I would say that claims of that nature would -- I guess might tend to make it more difficult to apply a complex-wide standard approach. And so it would fall into sort of the case selection portion of what's going through this process. you kind of have to -- I don't know that I can say universally, whoever says -- you know, makes that claim that we want to run through that process, I don't -- I won't stand here and make that claim, but I think it would affect -- you know, those kinds of issues would

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affect the case selection for what might go through
      this.
3 MR. GRIFFON: Okay.
4 DR. ZIEMER:
              Jim.
5 DR. MELIUS: Yeah. Just in response to what Tony was
      saying, I'm not -- what I was suggesting wasn't
      necessarily to add to the work for the Advisory Board,
      but in addition to the -- sort of the scientific
      confidence that we have in what NIOSH is doing, I think
      it's also our credibility issues and we have to
      remember that at the end of the process when we've gone
      through -- we're not going to be able to review
      everything through our contractor -- that what we don't
      review, the credibility of that has to be defended in
      some way.
6 Now if we say we've reviewed -- we may be confident by, you
      know, two percent or whatever it is of the cases,
      individual dose reconstruction reviewed that, you know,
      that's representative and that we've -- provides
      credibility to the process. But to some extent we have
      to think the same way about the site profiles, about
      these other Technical Basis Documents and I was just
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arguing for some sort of systematic approach and that -
      - and also to make sure that NIOSH wasn't expecting us
      to review -- to provide the technical review on all
      these procedures, even though it may be useful from a -
      - from NIOSH's point of view from -- in terms of the
      credibility of the application of these -- these
      processes, so -- I think it's just sort of coming to
      grips with that.
                          Tony, another comment?
9 DR. ZIEMER:
              Thank you.
0 DR. ANDRADE: Just a quick question for Mark. Do you have
      any idea on the percentage of these -- of the work in
      which we're -- in which our subcontractor's actually
      going to be looking at site profiles?
4 MR. GRIFFON: I don't recall off-hand. I mean the last task
      we laid out a pretty aggressive -- I forget the
      numbers, but we have a fair -- fairly high percentage
      of the overall site profiles. I think it was four AWEs
      and eight --
              I don't remember the numbers, but in terms of
9 DR. ZIEMER:
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      percent -- percent-wise, it's much higher than for
      individual cases. But that's understandable in terms
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      of the fact that many -- most of the cases come from a
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relatively small number of sites, actually.
2 MR. GRIFFON: Right.
3 DR. ANDRADE: Good.
                      Thank you.
4 DR. ZIEMER: Other comments or questions? Okay, thank you
      very much.
                  Thank you, Stuart, for that update.
                          IMBA UPDATE
7 Now we're going to turn our attention to the internal dose
      issues or the -- the calculation of internal dose.
      terminology is Integrated Modules for Bioassay
      Analysis, and we're going to get an update on that from
      David Allen of NIOSH. David.
2 MR. ALLEN: Okay. Can you hear me? All right. Thank you.
3 As Dr. Ziemer said, my name's Dave Allen. You've seen me
      before. It's been a while, but you have seen me
      before. And he mentioned I'm giving a presentation
      today on IMBA. And IMBA is the computer software that
      we've been using for internal dosimetry calculations.
8 As Dr. Ziemer already mentioned -- as Dr. Ziemer already
      mentioned, that stands for Integrated Modules for
      Bioassay Analysis. The difference between IMBA and the
      bulk of the commercially-available software for
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      internal dosimetry right now is that IMBA uses the
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current ICRP models. Any -- most of the other
      commercially-available software uses ICRP-30 models,
      which are a generation back.
4 I think it was easiest first to start with a little bit of
5
      history on the IMBA program, and up until the early
      90's, ICRP-30 was the current ICRP models for internal
      dosimetry. In 1994 ICRP published a new lung model for
      internal dosimetry, and that was the beginning of
      various new models, including biokinetic models. As --
      this particular lung model was considerably more
      complicated than the last, and as such, while they were
      producing this model, they also produced some computer
      programs to help evaluate that model. Once the model
      was published in 1994, the people that put that
      computer program together, NRPB, went ahead and
      packaged it as a software -- they connected --
7 DR. ZIEMER:
              Identify for everyone -- I -- I'm assuming, but
      it's probably not true, that ICRP is known, but maybe
      you should identify all the acronyms as you go.
0 MR. ALLEN: Okay, there's a lot of them.
1 DR. ZIEMER: If you remember them all.
2 MR. ALLEN: ICRP is the International Commission on
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Radiological Protection. It's basically the worldwide
      expert on internal dosimetry, recognized expert on
      internal dosimetry, as well -- radiological protection
      in general.
5 NRPB is the National Radiological Protection Board, which is
      I think semi-private/semi-government agency of Great
                Some of the people in that organization were
      Britain.
      involved when the ICRP committee for developing that
      lung model, and they developed computer software to
      evaluate that while it was being produced.
1 Once the lung model was produced, they connected it to the
      ICRP-30 biokinetic models, which were all they had at
2
      the time, and packaged that software in a form known as
              LUDEP then is like a hybrid of two different --
      LUDEP.
      your current model/old model type of thing. It was a
      DOS-based program. It was kind of clunky to run, but
      it was something that you had.
8 Shortly after the lung model, ICRP then began producing new
      biokinetic models, and as part of that, these
      individuals at NRPB were also involved with that.
      as new models came out, they produced new computational
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      models or modules that would do those calculations.
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And eventually all this was put together -- all these
      individual computational modules were put together into
      a -- one computer program, and that was known as IMBA,
      hence the Integrated part of the acronym -- Integrated
      Modules.
6 The first version of IMBA that I know of was IMBA-URAN, and
      after they -- NRPB copyrighted this -- these
      computational modules, they began putting IMBA together
      in an integrated fashion. IMBA-URAN, they were
      contracted to put together something a little more
      user-friendly for the CANDU reactors. That ended up
      doing only uranium, was all that would do.
      pretty limited in scope, but it did put everything
      together.
5 After that, near the same time, DOE contracted -- contracted
      them to put together an IMBA EXPERT version, which
      included more isotopes and a little more versatility --
      quite a bit more versatility. That took some time for
      them to complete, and then during that process, we
      contracted them to put together an IMBA-NIOSH version.
       That allowed for annual doses for a limited number of
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      isotopes, and gave us what we needed for this program
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on a limited basis. Once the IMBA EXPERT was done for
      DOE, we then asked for a modification of the software
      which included all the functionality of the IMBA EXPERT
      version, all those isotopes plus some additional
      isotopes, and that was all put together into what is
      now known as the IMBA EXPERT OCAS-edition.
7 Some of the features of the IMBA EXPERT edition is -- we can
      do up to ten individual intake regimes. By intake
      regime, that's -- that's a term given in IMBA, but it
      essentially is specifying the dates; the route of
      entry, whether inhalation, injection, et cetera; and
      also whether it's a chronic versus acute.
      together is one intake regime. You can specify up to
      ten intake regimes, so you can give somebody an acute
      intake on one day, followed by a chronic intake during
      another period of time, followed by an acute ingestion
      some other time, put it all together in one shot.
      it's good to have a number of those when you're talking
      about a career dose.
0 As I already mentioned, it can do inhalation, injection and
      ingestion. It can do -- whether -- it can do a chronic
      versus acute on any of those intake routes.
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solubility parameters can be specified as ICRP default
      solubility types, or you can specify individual
      parameters if you know more about the material the
      person inhaled or ingested.
5 Bioassay is -- we can use whole body counts, we can use lung
      counts, urinalysis or fecal sampling. There's also a
      few other bioassay types for specific isotopes, such as
      we can use thyroid counts for iodine exposures if -- if
      we have that data, we can use that then to determine
      intakes.
1 IMBA can be used to calculate the intake from that bioassay
      samples. It can be used to calculate dose from a given
2
      intake or the calculated intake. The dose can be
3
      effective dose, which is essentially a whole body dose,
      or it can calculate tissue dose -- tissue or organ.
      The dose that it calculates can be specified either 50-
      year committed or, more important for us, they can be
      specified as annual doses.
9 I'm going to take you through screen shots of the program.
0
      This -- there's not a good way to do this presentation.
       I'm going to take you through some screen shots just
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      because I think it will be a little quicker than
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letting you watch a computer program work while it's
      trying to work on the screen.
3 There's a number of pop-up screens and menus in IMBA.
      -- gets to be a fairly complicated program, but the
      three primary screens are the -- what I call the main
               That's the screen you get when you first turn
      screen.
      it on.
              The other two main screens are -- or primary
      screens are the dose calculation screen and the
      bioassay screen.
O That's the main screen. When you first turn the program on,
      that's what you see. For the most part, this is
      somewhat administrative data for the intake.
                                                    It will -
      - over in this side here -- it's divided into somewhat
      four sections. Starting over here, you're allowed to
      specify the particular radioisotopes you're interested
      in, and you can specify the exact intake if that's
      where you want to start. If you want to start from
      bioassay and calculate intake, obviously you leave that
      blank and that would come about later. The bottom
      section here is simply two buttons to take you to the
      other two primary screens.
2 And the top section right here, I have a little blow-up of -
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- a little clearer -- and as I said, this is somewhat administrative detail for the intake, all the information you need to put in ahead of time to start off with. Off to this side, you have -- you can specify the units you want the dose in, whether you want to work in sieverts, rem, et cetera. You can specify the units you want the activity to be in, whether becquerels, dpm, that sort of thing. 9 And up here is time. You can specify whether you want to deal with dates or whether you want to deal with time since the intake, such as ten days, 100 days, et In general, when you're dealing with an individual's individual case, the dates are usually the best thing to use. If you're dealing with a programmatic issue, like you want to come up with an excretion curve, then the days are probably better, the time. 8 This date you see here is nothing more than a reference date. If you're going to use time and specify the time since a particular date, then that reference date'll be the main thing. In the case over here, you see a zero in that time box. That means in this case it's -- this

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one represents 1/1/1980. If I were to put a ten in
      there, it would represent 1/11/1980, that sort of
      thing. And the reason for the reference date instead
      of just time since intake is 'cause you can do more
      than one intake. Some of these intakes could be years
      apart, so you -- the program needs one reference date
      to work with.
8 I mentioned on the other side you could do up to ten intake
      regimes. That's specified in this area. As you add
      more intake regimes to it, you get more tabs right
             Each of these tabs, once you click on them, you
      get an identical screen below here, and on this screen
      you can specify the route; you can specify the mode,
3
      whether it's acute or chronic; and you can specify the
      start date. If you click -- if you click on the
      chronic button, you also get another box to show up
      there to -- for the end date. As I said, somewhat
      administrative detail, but that's the -- the important
      detail, of course.
O DR. ZIEMER: Where does the solubility -- does that show up
      later?
2 MR. ALLEN:
             Yes.
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1 DR. ZIEMER:
              Okay.
2 MR. ALLEN: In fact, back to the main screen, the last
      section is down here where it says model parameters.
      There's a number of model parameters, and some of those
      will appear or disappear depending on the route of
      entry that you specified. Each of those allows you to
      change the parameters associated with that type of --
      those parameters.
9 I'm going to give you one screen shot -- that's the blow-up
      of the solubility. When you click on one of those
      buttons at the bottom of the main screen, you get a
      pop-up screen. In this case I clicked on absorption.
                          It shows you a pictorial
      I get this screen.
      representation of the model, as far as the solubility
      part of it goes. It shows you the actual values that
      are going to be used, and it also has buttons to allow
      you to pick the default solubility types in this case -
      - you know, F, M or S. You could also click on user-
      defined and then put in your own values here.
O DR. ZIEMER: What about particle sizes for inhalation, is
      that --
2 MR. ALLEN: That is one of those other many buttons on the
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model parameters, and I don't have a screen shot of I didn't want to come up with all of them. And that. that again is either default -- all the other models have ICRP default or user-input, user-specified. can see on this screen you also get an F1 value, which it could be specified on yet another screen, too. They're linked. And the help button will actually give you what the F1 values are for that particular isotope you've already selected to get to this point, and it'll tell you the chemical compounds associated with that -what that's -- the default solubility type are for that chemical compound, what the F1 value is for that default solubility type and the ICRP reference where that value came from. And you can simply click on that help menu, click OK and it'll put the value in there for you. 7 Okay. That was the main screen, and as I mentioned, there's two more primary screens associated with the program. It's somewhat backwards to do the dose one first, but it's the easiest one to deal with and then I'll get into the bioassay one. 2 When you click the dose button at the bottom of the main

values.

screen, you'll come up with the dose calculation screen. From this screen, you can start off by seeing off to the side -- it'll tell you the quantity -- the intake of -- the individual intake regimes that you're about to calculate dose for. In this case I've got three in there, and I think two of them are zeroes, so it's probably a bad screen shot to put up there, but that's what I ended up putting up. 9 Over here you simply have a calculate button. Once you hit calculate, this will give you the effective dose from this -- these intakes that you have specified -- these ones that are listed over here. It'll also give you the organ dose for each intake regime, as well as the total of all three. And this screen you can see -there's a slide bar. There's a lot more organs than there's -- you can see from this screen, but... 7 These that you're seeing here, the effective dose and the organ doses, are all 50-year committed doses. For purposes of our program, that's not very useful. don't use that much, but it is there and it's a great QA for the program itself to verify it against the ICRP

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1 Up here in the corner you see a button that says calendar.
      What that button allows you to do then is get to where
      you can calculate the annual doses. You get another
      pop-up screen.
5 This is the annual dose calculation screen, and what it
      allows you to do is input the start year.
                                                  There's a
      pick list of 30-some organs that you can choose from.
      There is the end date, which would be the date of
      diagnosis for us. And once you have all that
      information entered in, you hit start calculate.
      takes a little bit of time, but it'll run through the
      calculation and it'll give you the annual dose each
      year from the start year to the diagnosis date. And
      you can see from the screen, the last year it's going
      to be a partial year, from the beginning of that
      calendar year to the date of diagnosis.
7 The last two buttons are down here. That allows this
      information to be copied to the clipboard or to be
      exported as an ASCII file, which makes it much easier
      to use. You can simply copy it to the clipboard, paste
      it into say Excel if you wanted to, and then you have
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      all the values you want to...
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1 Okay, the last primary screen is the bioassay primary
      screen, and right here are two tabs. These are keys to
      the screen. What you can see on there right now it
      says bioassay to intake, and that would allow you to
      calculate an intake from the bioassay. If you click
      the other button, the other tab here, it gives you a
      little different information here. It allows you to
      predict the bioassay measurements from a given intake.
       Sounds like a subtle difference, but it's an important
      difference.
1 In this case I have urine selected here. I'm on the
      bioassay to intake screen. When I hit calculate, it's
      going to take the data that I've put in there, the
      measurement data that I've put in there, try to fit the
      data the best it can to all the intake regimes that I
      put up there as far as the intake dates, routes of
      entry, all that. It'll try to fit that data the best
      it can to that and come up with the intakes, the actual
      quantity of -- that was the intake.
0 If I go this button here, I have to specify dates that I
      want it to calculate or predict what the bioassay would
           I click a calculate button on that screen.
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doesn't change the intakes at all. It simply
      calculates what a bioassay would be from those given
      intakes.
4 Over here on this side I have actually three identical
5
      areas, though they don't look identical right now.
      Each of these areas allows you to choose a bioassay
      from a pick list -- bioassay types, such as urine,
      fecal, et cetera. It allows you to have the table or
      the graph, and each one is identical, so in this case
      I've got the table data for urine and a graph for the
      urine samples. I can see them at the same time.
      makes life a little easier when you're working around
      with the data. If I also had lung count data, I could
3
      put that up there, either the table or the graph.
      had a number of things I could put all three graphs up
      there, however I wanted to work it out.
7 Each of these screens have a little tool button.
      click that, you get a more detailed screen or a bigger
      -- full-page screen of those individual spots.
      just a little blow-up of the exact same thing I just
      showed you. I thought it might be a little easier to
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      see.
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1 There's the full-screen view of the table data, and you can see you have two different colors -- color codings. The green is for the prediction. That's if you want to go from the intake to the -- predict what the bioassay would be. The input in this case would be the dates that you're interested in. In the case of urine also, the collection period. 8 The blue area is actual measured data. It would be a date that somebody was sampled, the collection period and the actual measured quantity. The data type, there's actually three options here. You can specify whether that's a real quantity, an actually measured quantity, or if it's simply a less-than. The less-than LOD stands for less than limit of detection. 5 The third quantity I don't have on there is simply excluded. From time to time you might get an outlier that doesn't seem to make any sense, and that would allow the dosimetrist to select excluded for that dataset, and at that point the program will ignore it. It'll still plot it on the graph, but that's the only thing it'll do with it, so if you can ignore what looks like 2 an outlier and then all of a sudden everything fits

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real well with the models, then you've got a chance
      that it's -- it really was an outlier.
3 The other two are semi-self-explanatory. They show you the
      measurement error and the error distribution. Your
      options there are normal or lognormal.
6 It's a little better blow-up of the graph. When you click
      the tool button on a graph, then you get this blow-up.
       I've cropped a little bit of it out so you can't
      really see what's going on down there, but that is
      essentially a lot of administrative type information --
      what scale you want, format for data, how many decimal
      places, that sort of thing. As far as the scale, what
      you see here are the days since that reference date
      back on the first page. Off to the side here is the
      actual measured -- or the quantity that you're trying
                   In this case I think it was in picocuries
      to measure.
      per day for urine samples, I believe it was.
8 The blue dots are the measured values, complete with the
      error bars.
                   The black line that you see here is the
      fit that IMBA did to that -- that data, based on the
      intake regimes that you gave it. The green -- I don't
2
      know how well you can see that -- actually follows all
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2

the way along this black line, and the green is the predicted bioassay. The reason you see such a difference here is the graphing function itself is not that sophisticated, and it's simply connecting dot to dot that the program has calculated. When you see something like this with the black line and it's say several weeks after initial intake, someone might be thinking it should be higher and they could be coming down by then. What the predicted bioassay allows you to do is select some dates in between there. is relying on the dates that the sample was done. The green, you can select dates to see what the model predicts to see if it is realistic. In this case, in between there you can see where the urine should have jumped way up and then been coming down by the time that first sample was taken, after -- so you get the idea of what you're saying from this intake regime, what you're predicting.

9 The last thing I wanted to show you was I got a split screen here of -- back to that bioassay screen, and I wanted to show you the utility of graph and why a picture's worth 1,000 words. Off to the side here, this is a

graph of urinalysis for a particular individual uranium exposure over -- over time, various different intake regimes. A number of intake regimes were put in, IMBA calculated the values for the urinalysis and this is what he -- the black line versus the blue dots or how it fit together. It seems fairly reasonable on this. 7 Down below here the blue dots are lung counts that were done The intakes were not determined from 8 on the person. lung counts. They were determined in this case from urinalysis, and then the lung count data was predicted, and that's the green line you see up here. And you can see the green line doesn't seem to match the blue lines very well at all. It seems to be considerably higher. That pretty much tells the dosimetrist, you know, some of the assumptions are wrong somewhere. In this particular case, this was assumed to be a type S material, and IMBA then allows you to go back, change the solubility assumption to, in this case, type M, redo the same thing, and you can see there that you can not really tell any difference in the fit. You can fit urinalysis data very well with type S or type M, 2 different quantities, but the intake regime was the

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same.
2 The lung count data, on the other hand, when you compared,
      all of a sudden that green line fits right through all
      the blue dots. It seems to be most realistic in this
      case to be type M type material.
6 And that's pretty much it. I'm looking at a lot of stone
      faces and no questions. I know it's a dry topic --
8 DR. ZIEMER: I'm trying to see where the -- where are the
      data points in those two lower curves? Are they down
      near the axis? Is that what --
1 MR. ALLEN: Yeah, there's a row of data points right there -
3 DR. ZIEMER: Okay, I see.
4 MR. ALLEN: I put the axis in that situation -- I mean I
      could have spread it out a little more, but I put the
      axis so I could get this green line on the screen.
                                                          And
      then in this case, I just wanted to compare apples to
               I left the axes the same.
      apples.
9 DR. ZIEMER: Okay, thank you very much. Let's open the
      floor for questions. Tony.
1 DR. ANDRADE: Dave, you mentioned in one of your screens
      there that either the model predicted points below the
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detection limit or data were entered that were below
      the detection limit. Which was which and can you
      explain that to me again? That one kind of took me by
      surprise.
5 MR. ALLEN: Went by a little fast?
6 DR. ANDRADE: Yeah.
7 MR. ALLEN: That was measured values below the detection
      limit, so --
9 DR. ANDRADE: Measured values?
0 MR. ALLEN: In other words -- example, somebody got a
      urinalysis and the results were less than one picocurie
      per day or some value --
3 DR. ANDRADE: Right.
4 MR. ALLEN: -- you can put in that value and say it was less
      than.
             That's what the program allows you to do.
6 DR. ANDRADE: But wouldn't a reasonable health physicist
      have a decision limit that's greater-than? I mean
      using classical statistics, okay, not Baysian, but
      classical, wouldn't you have a detection limit that's
      certainly somewhere way above your -- I mean a decision
      limit way above your detection limit?
2 MR. ALLEN: Yes, you would. Unfortunately, a lot of cases
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we have, there's going to be no detectable samples. We're stuck with a lot of less-thans. This at least allows us to plot them out and saying they're lessthan. If -- if there's enough detectable sample that we want to ignore what was less-than, we can exclude those with the exclude type and see how everything fit with what was actually detectable and make sure that's reasonable for that situation. There's a number of options that are available there.

0 DR. ANDRADE: Okay. So you're artificially establishing a floor that's actually below your detection limit for your system.

3 MR. ALLEN: No, actually what it's doing is it is using a maximum likelihood method on the fit and by establishing it -- the value is less than detection limit, it says it has to be in that range between -somewhere in that range between zero and the detection That way it tries not to fit it. If it's going to come out way above that, it's not going to try to predict that intake. It'll predict one that's going to put it down below there, but it doesn't give a lot of weight to it, just somewhere in that range.

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1 DR. ANDRADE: Thank you.
2 DR. ZIEMER: Jim and Gen.
3 DR. MELIUS: Gen was first.
4 DR. ROESSLER: I have a question with regard to the IMBA
      history. You mentioned that NRPB developed these
      biokinetic models using the new ICRP models in about
      1994?
8 MR. ALLEN: Well, 1994 was the lung model published -- when
      the lung model was published. NRPB did play around
      with some computer models before that, and that
      accounts for some of the values in that publication.
      Those for the most part came from Allen Birchell*, who
      works for NRPB.
4 DR. ROESSLER: And then you mentioned that the IMBA models
      or model was developed. Was that done by NRPB?
6 MR. ALLEN: Yeah. NRPB copyrighted the calculational model
      that they use for the LUDEP and the lung model.
8 DR. ROESSLER: And then -- and when -- when was that?
9 MR. ALLEN: The copyrights -- the individual models I
      believe were copyrighted at various times.
      actually several modules, one of which is a lung
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deposition*, one is a --

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1 DR. ROESSLER: It's more recently, though, I guess?
2 MR. ALLEN: Excuse me?
3 DR. ROESSLER:
               More recently?
4 MR. ALLEN: Probably post-'94 when it was copyrighted. And
      then as the biokinetic models were developed, they
      copyrighted new modules.
7 DR. ROESSLER: Up to this time, up to your presentation, I
      was thinking that NIOSH had developed IMBA.
      you're really doing, you're using the program developed
      by NRPB.
1 MR. ALLEN: Right, what it amounts to is copyrighted
      software -- or copyrighted calculational engines, and
      we asked for the front end -- everything you see pretty
      much, we asked for that to be developed, a user
      interface.
6 DR. ROESSLER: My real question is who validated the model?
       Did NIOSH do anything -- I mean I want to know if it's
      working right, and I know NRPB probably did, but I'm
      wondering what you did to -- to validate it to make
      sure that that's the model you wanted to use and that
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2 MR. ALLEN: Right. NRPB did a lot of V&B* on it, quality

you're getting the right answers.

control, every time they develop a new model on there, they do a lot of quality control. Also they were pretty much the only thing that had any sort of credibility or V&B out there at the time we were looking for something. And when we get in a new version, what we've done is used the committed dose section of that. We can put in -- at one becquerel intake, use the committed dose and see if we get the right effective and the right organ doses according to NRPB publications, and that's what we're claiming to use is NRCP -- NCRP models -- I'll get it right -- ICRP models. We're claiming to use the ICRP models. program with the particular input gives us the right output that we can tell from those publications, and that's -- that and maybe a little bit of more validations is about all we manage to do in-house, but we have the NRPB quality control documents, also.

8 DR. ZIEMER: Jim, then Mark.

2

9 DR. MELIUS: Well, actually Gen asked one of my questions, though I'm still a little bit concerned with just the -- the validation issue. So -- so how do we know that this is giving you the right -- the correct answers for

other situations? I mean it -- assume there's been more to the quality control that went into developing this than -- than what you described. 4 MR. ALLEN: NRPB has put a lot into it. What we've done in-5 house is -- first thing is to -- for each isotope, we put in one becquerel and see if the effective dose is what's -- matches what's in the publications. Also see that the committed dose to organs matches what's in the publications. After that, we've determined -- we calculated annual dose for 50 years, put them in Excel, added them up and made sure that matched. So it's -you know, annual doses at least match -- or add up to the 50-year committed dose. 4 We've also -- using ICRP-78 where we can with the dates that are in 78 for bioassay, we can predict bioassay -- put in, you know, standard input, like a one becquerel intake and predict what the bioassay should be at fiveday, ten-day, 100-day, whatever's in the ICRP publication, and we can verify that that matches with the publication. And we've done a little bit of work matching up with -- Potter published a whole magazine 2 of tables for bioassay analysis and we've done some

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spot-checking against that.
2 DR. MELIUS: Secondly is the issue of how you use it, and I
      guess I'm a little -- well, I don't -- confused or
      concerned, but at one point you mentioned that you're
      going through and when you find an outlier, you exclude
      it.
7 MR. ALLEN: The option's there for the dosimetrist.
8 DR. MELIUS: Okay. But is that what you -- you're actually
      doing? 'Cause I mean I --
0 MR. ALLEN: It can be. I mean errant bioassay samples do
      happen.
               There's not unusual to even get a number of
      samples, you can watch urinalysis coming down from an
      acute intake and then one of them's zero.
4 DR. MELIUS: Yeah.
5 MR. ALLEN: The next one seems to be following the curve.
      That zero, if you assume it's real and the computer's
      looking at it, it's going to drive that intake down to
      try to fit the data, so the dosimetrist is allowed to
      exclude that and see if the data fits better without
      that anomaly.
1 DR. MELIUS: Yeah, but then what -- how do you take that
      into account when you're doing your actual dose
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calculation? How do we know that your assumptions -your assum-- that you're excluding it 'cause it's a bad bioassay sample versus that you're making the wrong assumptions because of the -- you have poor information? And then how does that get into -eventually get into the IREP model as -- in terms of your certainty about that dose? I guess -- and I'm going for -- trying to get at how much, you know, individual, you know, (Inaudible), if ten different health physicists used the same data, would they all come up with the same, you know, calculation or -yeah, yeah, that's --3 MR. ALLEN: For internal dosimetry, no. You will get ten different answers. Hopefully they'll get the -- our -our job is to make sure we get the same side of 50 percent probability. The dose should be reasonably close, the intake. 8 For the most part, as far as the uncertainty, at this point we can -- I can answer that that we've avoided it. the most part we have tried to overestimate or underestimate bioassay. When we have data on a curve 2 or data showing on a curve that's considerably higher

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or considerably lower than that data point's indicating
that it's definitely an overestimate or definitely an
underestimate, therefore we can bound the intake that
way. And that's probably going to be a big portion of
the cases. We can bound it that way and not deal with
the -- you know, whether it's ten percent error or 50
percent error or whatever.
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8 DR. MELIUS: No, I -- I understand that part. I just get
9 concerned about the ones that you can't do that on and
10 where your assumptions are going to be sort of critical
1 to the outcome of that case, and sort of how do we get
1 consistency in doing it, I think is the -- is the
1 issue. I think Tony has a...

4 DR. ZIEMER: Mark and then Tony.

in this newest version, what are the radionuclides
available now? I know in -- I saw a recent version and
it had just limited radionuclides built in and I wonder
if -- if you've got all the radionuclides you need for
this program available now.

1 MR. ALLEN: We have about -- I believe the number's 54 now
2 in there. It's all the important ones you would want

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with DOE complex and probably not every single one.
      There's always an odd case out there, especially at a
      national lab. For the most part, what's missing either
      has a short half-life or a short biological half-life,
      and we can use the 50-year committed published doses --
      the person got it all in the first year and we can
      simply use that without the computer program.
8 DR. ZIEMER:
              Tony?
9 DR. ANDRADE: Dave, certainly within the laboratories we
      have ways of ensuring that the spikes and the zeroes
      are true or not true. We run blanks and we also run
      spikes, along with the real bioassay samples. And that
      is true for both alpha spectroscopy as well as for mass
      spectrometry. And I guess my question to you is do we
      pass along or do you use all of the raw data that we
      collect? Or do you use our final values?
7 MR. ALLEN: We ask for the raw data and that's what we use.
8 DR. ANDRADE:
               Okay.
9 MR. ALLEN: If we can -- all the way down to counts per
      minute if that's what we can get, but that's pretty
      rare to get ahold of that. That's generally going to
2
      be a urinalysis result in say dpm per day or whatever
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the appropriate unit is for that particular isotope, and that's what we start with. 3 DR. ANDRADE: Okay. 4 DR. MELIUS: Yeah, the issue's been brought up before here about the accessibility of this software for people that are not directly involved in the program, and I may be wrong, but I thought Larry or somebody was going to look into that issue. Am I remembering wrong? don't want to put you on the spot, but -- or have we thought about that more or -- that whole issue? 1 MR. ELLIOTT: We've already answered that. 2 MR. ALLEN: I've encouraged the vendor to make a publiclyavailable version that he can sell. They are looking 3 at that. They have not made one yet. There is a version they're putting together called IMBA Professional, and he's trying to put together a light version -- is what he calls it, IMBA Professional Light -- that has much fewer functions that might be -possibly be affordable is what he's shooting at. is not available yet. He is -- they're -- haven't sorted that out, but individuals would have to buy or 2 organization or whatever, 'cause there's licensing

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issues with the copyrighted software.
2 MR. GRIFFON: And how -- just to follow-up on that, how
      about availability to the Board or to the
      subcontractor, can --
5 MR. ELLIOTT: You -- the Board and your contractor, as well
      as our contractor, all have access to the IMBA-NIOSH --
6
      or IMBA-OCAS as special government employees or as
      contractors to the government on the program. But Dave
      is actually right and accurate in his statement.
                                                          The
      ICRP models and the calculation engine from NPRB (sic)
      are copyrighted and protected and we can't distribute
      those to the public without a user's license, and
      that's the issue. Somebody has to pay for that.
4 MR. GRIFFON: And you say they're available to the Board.
      mean can we -- can we physically get a copy sent prob--
      before the next meeting or how can we move forward on
      this? It'd like to get a disk copy sooner than later.
             It can be done. We've got -- I don't know how
8 MR. ALLEN:
      we'll work out the details on -- the licensing
      agreement allows NIOSH to use it and any of its
      subcontractors for the purposes of the OCAS --
2 MR. ELLIOTT: Am I correct, it's not in a CD form, though,
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it's in a -- it goes on a server. Right?
2 MR. ALLEN: No, it's in a CD.
3 MR. ELLIOTT:
              It is in a CD.
4 MR. ALLEN:
             Yeah.
5 MR. ELLIOTT:
               Okay. Well, we'll get that --
6 DR. ZIEMER:
              Okay, you can work with them then and --
7 MR. GRIFFON:
               Okay.
8 DR. ZIEMER: Yeah, Mike has a question here.
9 MR. GIBSON: Can you tell me -- do you know off-hand how
      much of the data you get from these DOE sites is
      assumptions, default factors, solubility class, the
      date of intake?
3 MR. ALLEN: For the most part, if it's bioassay -- that's
      why we want the raw data. If it's bioassay analysis,
      it's a mass spec or an alpha spec or even a gross
      alpha, that's -- counts or activity in that urine, for
      example. There's no assumptions that went into it.
      It's a laboratory analysis. And then from there we
      have to make the assumptions as far as solubility and
      all that. And the assumptions we use are the reason we
      have to put together these site profiles that was the
2
      last lecture. We have to have some idea of what
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material they had so that we know what type of
      solubilities that would be associated with it, and then
      we make sure that the bioassay actually fits that data,
      if the assumptions are accurate or not.
5 MR. GIBSON:
             But your -- are you looking at the date of the
      bioassay -- was taken or --
7 MR. ALLEN: Yes. Yeah, we -- we get the date that a sample
      was taken; you know, what was taken, such as a urine
      sample; how much, was it a 24-hour sample or was it a -
      - you know, an allotment; the actual results, such as a
      gross alpha or it could be picocuries per liter of
      plutonium, for example; what isotope if they have that.
      Everything we can get, we get. We're not shy about
      asking for it.
5 DR. ZIEMER: Okay on that, Mike, or did you have a follow-
      up?
7 MR. GIBSON:
              I don't know if I understand it all, but I --
      that'll answer for now, yeah.
9 DR. ZIEMER:
              And you're saying basically you're not
      utilizing assumptions that may have been made on the
      site 'cause on the site they also presumably do some
      sort of dose calculation, in many cases the 50-year
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committed dose --
2 MR. ALLEN: Using ICRP-30 models --
3 DR. ZIEMER:
             Right.
4 MR. ALLEN: -- so what they've used -- the doses they've
      calculated are not necessarily good for us, and very
      few sites have ever calculated an annual organ dose.
              So you don't find there's any value in looking
7 DR. ZIEMER:
      at what they may have ultimately calculated for tissue
      dose or organ dose?
O MR. ALLEN:
             There could be some value in it, especially -- I
      mean information's always limiting, and if that's all I
      can get is a calculated dose, you can pretty much -- if
      you get enough details, you can back-calculate what the
3
      bioassay was that that came from. Also as far as the
      solubilities, same general -- you know, we hope that
      we're not complete ends of the spectrum on what they're
      -- have been assuming and what we're going to assume.
      There should be some reason if there's a -- if we're
      completely different.
O DR. ZIEMER: He's got a follow-up.
1 MR. GIBSON: If you had like a super-Y class of plutonium,
      how would you be able to distinguish that or could that
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mask your raw data out of the bioassay...
2 MR. ALLEN: It really couldn't mask the raw data, but it
      could mask the dose and the intake that you're
      calculating from the raw data. And the way that can be
      handled -- as I said, we have default classes that can
      be picked, but we can also input our own user input.
      For us to use a super class-Y plutonium, we would
      probably put together a Technical Information Bulletin
      evaluating a particular site, what they had, saying the
      solubility doesn't follow the defaults and we have more
      information, and in that case use these absorption
      parameters for this site, is how we would handle that,
      and we haven't done that yet. What's -- we haven't
      changed from any defaults yet, but we're still kind of
      young into that part.
6 DR. MELIUS: Just one -- I think it's a brief question, but
      back just to the validation. Have you documented the
      validation you've done?
9 MR. ALLEN: Part of our contract for the upgrade is a whole
      documentation on all the V&B that NRPB has done.
1 DR. MELIUS: Okay. And then your further --
2 MR. ALLEN: Our further evaluation, we have documented not
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in a very formal manner, okay? That's one of those
      things you just never seem to get to. We've got all
      the numbers and it's a matter of writing down something
      and documenting it.
5 DR. MELIUS: I just think that would be helpful to do.
      know it's hard to get to, but it's one of those things
      -- things that I think at some point, if questions are
      raised, it would be good to have.
9 MR. ALLEN: Yeah, I understand. It's just once you get --
      once you have run the numbers, you know it works, the
      actual --
2 DR. MELIUS: No, I -- I --
3 MR. ALLEN: It tends to get pushed to the back burner.
4 MR. GRIFFON: Yeah, just one final thing on the validation
      side of it, do you know if there are any plans to
      update the CINDY* code to be ICRP-60/66 compatible? I
      know right now it runs in 30, and if that's going on,
      that may be another tool that you can validate against
      or whatever. I don't know if that's happening.
0 MR. ALLEN: I don't know if that's happening. I haven't
      heard of that. I know there is a number of -- a number
      of other codes out there. For the most part, they're
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kind of home-brewed. Like Potter put one together, and
      there's someone else that put a math CAD-1 together, I
      think, and --
4 MR. GRIFFON: French or --
5 MR. ALLEN: Nothing that's very versatile for what we're
      doing.
7 DR. ZIEMER:
              Tony.
8 DR. ANDRADE: A quick answer to Mark's question. When we
      were asked by DOE to assist in the development of IMBA,
      we were all forced to shell out some big bucks, and so
      in doing so DOE elected for us to invest in this
      particular code. So if CINDY is being upgraded, then
      it's got to be getting done sort of at a -- at the
      grassroots level somewhere.
5 UNIDENTIFIED: Good answer.
               Just -- just one final question, Paul.
6 MR. GRIFFON:
      curious if you've -- just in individual cases, you
      mentioned bioassay data a lot. Have you had the
      occasion to use air sampling data to validate your dose
      calculations from your bioassay, and I don't know how
      often you're able to get the air sampling data that
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might be appropriate for certain individuals, but have

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you had the -- have you done that frequently or --2 MR. ALLEN: No. I would love to, but getting the air sample data, especially from like a major DOE facility and 3 correlating that to an individual throughout a 20-year career is virtually impossible. We can get some ball park estimates if we could get the data, but because it wouldn't be that useful in that situation, we haven't gone -- trying to get it. It's also fairly difficult to get ahold of, as far as 20 years, 30 years back. You can get some general ideas, but the details is hard to --

2 MR. GRIFFON: I'm not sure I agree with the it wouldn't be very useful part of that statement, but otherwise I agree with you. I mean -- I mean I think there might be some usefulness for certain priority operations or areas within certain sites to have that as a backdrop, and if you knew a person worked in that facility over a certain period of time, you could do some cross-checks or -- and I certainly have noticed also that that's lacking in the site profile documentation, too, so I would encour -- I think that's a useful tool to -- if nothing el-- I mean I know it's probably going to

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increase your uncertainty in your overall estimate in
      many cases, but it's another piece of information, as
      you said earlier, which I -- you know, it may be
      valuable in certain circumstances.
5 DR. ZIEMER:
              Thank you. We're at the lunch hour.
                                                    I think
      it's appropriate now for us to recess for lunch and
      we'll come back together at 1:30.
8 (Whereupon, a luncheon recess was taken.)
        WORKGROUP ON OPTIONS FOR EVALUATING INTERVIEWS
O DR. ZIEMER:
              Thank you. We're ready to call the
      meeting back to order. I trust you all had a good
      lunch and are ready for another working session.
3 We're going to begin our afternoon session with a working
      report from our workgroup on options for evaluating
      interviews. Dr. Melius has been the Chairperson of
      that workgroup and he's going to report to us and
      perhaps make a recommendation.
                            (Pause)
9 Okay, Dr. Melius.
O DR. MELIUS: Okay. Since our last meeting in St. Louis, the
      workgroup has had one additional conference call -- was
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it last week? Yeah, last week, Wednesday. We met by

conference call and discussed -- and had received some additional information from NIOSH, which were the -basically the ORAU procedures for doing the interviews and scheduling the interviews and so forth. the information that we had received and I think recognizing that -- that NIOSH's and ORAU's program to sort of review the interviews and some of the quality assurance/quality control measures were a work in progress -- they were developing these really as -- as we were meeting, and as the program was getting -getting implemented -- we've -- came up with a set of recommendations which I think everyone has in front of them here about things that might -- these covered two One is things that NIOSH might do as part of -areas. and actually it may currently be -- already be doing as part of its quality assurance/quality control program for the interviews -- that would be helpful for the Advisory Board if some of this -- these steps were -or procedures or events were captured in some way as part of a -- the database so that we would be able to -- the Board would be able to go back at some point in time and evaluate these or our contractor might as a

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way of evaluating the interview process in the context of the dose reconstructions that are going on. And so we've made a basic recommendation there to -- to NIOSH. It's not meant to be overly prescriptive for NIOSH, but to -- that basically the program be further developed and that -- and we've given some examples of things that might be captured. Almost all of these examples are things that NIOSH -- in fact all of them may be very well things that NIOSH or ORAU is already doing. And the question is just to make sure that there is some record that keeps track of these, and in some sense a tracking system is -- that would allow review.

4 The second part of our recommendation is that -- then -that as part of the dose reconstruction -- dose construction review process that would undertake that then as currently cons-- currently described, this -our dose reconstruction will be -- program review will also be evaluating the outputs from the interview, the way the interviews are recorded, and also some of the other information that's kept in the individual record to that. And so that's what's captured in this second

recommendation there. 2 Now I've circulated this to the members of the working group, also to Paul. Paul's listened in and participated in our last conference call. I really didn't receive any comments or corrections from the workgroup, but they're free to correct or whatever as we go along, but -- but I think it's more important sort of the concept -- again, to go back, what we're trying to deal with is the issue of should there be -should the Board be either repeating or taking some steps that would be more intrusive in terms of evaluating the interview process, be that a second interview, independent interview, a review of a transcript of an interview or recording of -- of interview and that the Board is probably split on that issue as to whether or not that should be done or whether that's too much of a intrusion or imposition on the people that -- on the claimants and so that these steps in place and based on the results of -- of this review, then the Board at a later point in time could make an assessment as to whether or not a more 2 intrusive form of review of the interviews might --

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would or would not be necessary.
2 I don't know if any other members of the working group have
      any comments you want to add to that.
4 DR. ZIEMER: Okay, before we take comments, the Chair is
      going to interpret this as a recommendation -- it is a
      recommendation from a working group and as such
      constitutes a formal motion before the Board, doesn't
      require a second. And so with that as background, we
      can have comments, which could include modification.
O Let me also add, and as Jim indicated, I did listen in on
      this and I want to make sure -- particularly that the
      NIOSH staff understands that this Board is not
      mandating specific things that NIOSH do. We do not
      want to micro-manage NIOSH. The -- as I understand the
      intention of item one and the list of (a) through (f)
      is that in fact these are the kinds of records we would
      like to be able to sample as a Board, and if they
      existed, we would then in turn be able to evaluate the
      -- the process, the interview process more readily.
      Whether -- I think we believe that probably most of
      these exist in some form -- either formally or
2
      informally -- but the whole idea there was to identify
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the kinds of things we -- that probably the Board would
      want to sample that would make it much -- make the
      audit more readily conductible. Is that a fair --
4 UNIDENTIFIED: And informative.
5 DR. ZIEMER:
              And informative. Is that a fair statement,
      Jim?
7 DR. MELIUS: Yeah, and -- and --
8 DR. ZIEMER: And let's begin with Wanda, and then we'll jump
      down to Tony. And you can speak for or against the
      motion or modify or just comment.
1 MS. MUNN:
           My apologies for not having gotten back to our --
      our working group chair with a couple of comments that
              I think they were both captured by comments
      I had.
3
      that we made during our actual discussion, but I felt
      perhaps were not fully gathered here. I had thought we
      might meet once more before we actually presented
      anything in writing to the group and -- but most of --
      most of what I had -- had -- most of the changes I had
      made were purely editorial. They didn't change the
      sense of what was going on here.
1 The one thing that I did not feel was captured that -- was
      the suggestion that I made, which remains important in
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my mind, that it would be most helpful from an auditing
      point of view to have a single document where a record
      had at least been signed off on by individuals who had
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      done these specific actions that were listed here.
      don't think that such a document would be an undue
      burden if it went along with the case file, wherever it
             But that's something that certainly would be
      simply a suggestion as a potential tool that might be
      considered. And I can see no reason why it would have
      to be written, necessarily.
                                   It's --
1 DR. ZIEMER:
              Wanda, are you suggesting a specific change?
      seem to recall you characterized that as a tracking
      system in the phone call. Was that --
4 MS. MUNN:
           Yes.
5 DR. ZIEMER: Am I thinking about the right thing?
6 MS. MUNN: Yes, you are. Yeah, I was thinking about a
      specific document that would serve as a tracking
      document.
9 DR. ZIEMER: Well, is that item (f) or is that different
      than item (f)?
1 MS. MUNN: Well, I think -- I interpreted item (f) to
      incorporate that, but perhaps -- if one had not heard
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the discussion, I thought just reading (f) as it was perhaps would not make it as clear as what I had in mind. I had a simple sheet of paper in mind which would be a check-off or a sign-off document for various steps that needed to be gone through, which were over and above the mechanical processes that are done electronically. But no, I'm not asking for any changes. I just wanted a clarification statement. 9 DR. ZIEMER: Thank you. Tony? 0 DR. ANDRADE: With respect to Wanda's comment, I would just like to say that if -- if an electronic system is put into place, such as we suggest here, then a paper which would essentially be a traveler, as we call it, could be generated from that electronic system's -- at any particular point in time, at -- with any particular case, such that the audit function would be very, very easy to accomplish. So I think that if we can accomplish what's written down in item (f), then the paper document would be a natural. It'd be -- it would follow on naturally. 1 Again, in keeping with what Paul said, I, too, do not want

to be overly prescriptive or to try to dictate the work

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that -- that NIOSH should do. However, I did want to
      point out that a QAQ (sic) system is -- is really meant
      for those people that are implementing these -- these
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      processes. It's meant for their own quality
      development and improvement. And therefore we should
      not lose sight of the fact that NIOSH should own the
      program, should evaluate the program. ORAU should be -
      - should use the procedures that are developed, and
      that the Board should also keep track of what's going
           And I believe this is also an item for our
      subcontractor to look at. Am I wrong?
2 DR. ZIEMER:
              There is part of the task which is --
3 DR. ANDRADE: There is part of a task there might --
4 DR. ZIEMER: -- this, and in fact when we're talking about
      doing this in terms of "us", that includes our
      contractor.
7 DR. ANDRADE: Right. Okay. I, too, wanted to hand over a
      couple of -- both editorial and maybe one or two
      substantive comments, and I don't know if this is the
      appropriate time to delve into those.
1 DR. ZIEMER:
              That's fine.
2 DR. ANDRADE: Okay. I try to skip over the editorial piece.
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Let me just get back to the very last paragraph in
      this draft document, and it refers to the fact that --
      I think next to the last -- sorry, the last sentence.
      However -- it reads: However, the need for this should
      be re-evaluated at a later time -- and we're talking
      about re-interviewing claimants -- based on the results
      of the dose reconstruction review and the
      implementation of the QA/QC program described above.
9 Well, QA/QC is meant for quality improvement, meaning
      improving processes into the future. They should not
      be looked at as an avenue to go back and look at things
      retrospectively. Which brings us to the heart of the
      matter.
4 I think that we as a Board should vote and should decide
      once and for all whether re-interviewing is actually
      even on the table. I believe it shouldn't be.
                                                      Ι
      believe it's onerous. I believe that the only people
      that are going to be called are those people who have
      had their claims rejected and that it's going to be
      just a heart-wrenching experience for those people.
      And I really don't see any tremendous incremental value
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      added in even thinking about that.
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1 So with that, you know, I would -- I would essentially vote
      for taking out most of that sentence, starting with the
      word "based". But of course that really can't be done
      until this Board comes to -- comes to grip with that
      issue, and I really do believe that we should do that.
6 DR. ZIEMER: Tony, I'm not sure whether you're making a
      motion to amend or simply at this time reflecting that
      viewpoint...
9 DR. ANDRADE: Well, I --
O DR. ZIEMER: Could you clarify for me?
1 DR. ANDRADE: Okay, Paul. I guess there'd be a two -- two-
      phased approach to this. One is, I think that the
      Board should discuss whether or not retrospective or
      re-interviewing is even on the table for us to
      consider. That's one.
6 And if it is not, then I would move to amend the draft as it
      -- as it -- as it stands, at least for the --
8 DR. ZIEMER: Yeah, I might point out that it -- it may turn
      out to be a moot point whether or not we include it in
      this document.
                      The Board could always -- even if this
      were deleted, the Board could at a later date decide,
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      for whatever reason it wished, that something different
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should be done. We're not binding ourselves in one direction or the other. I suspect that the statement as it stands simply points out that the door could still be open for that possibility in the future. You're suggesting let's not even open it --6 DR. ANDRADE: I don't think we --7 DR. ZIEMER: -- and I'm saying that even if we did that, there would be nothing to prevent the Board in the future from changing its mind in any event. And I say that in the context where I myself have been basically opposed to the idea of re-interviewing, if only for the fact that a re-interview is not in fact the same as the original interview. It is different in time and in pla-- space. The interviewer would be a different person than the original one, presumably. You could not reproduce the conditions of the original interview. You might in fact elicit different responses from interviewees. You might elicit things that the interviewee did not even think of the first time around, so it's very difficult for me to imagine a reinterview as a quality check on the original interview 2 so much as these items, which are a way of getting at

- the issue of whether or not the information from the interview was properly captured and used in the record and in the determination of the eligibility of the person for compensation.
- 5 But be that as it may, I think we need to hear from others.
- I guess you're not making the motion at this time, or are you?
- 8 DR. ANDRADE: No.
- 9 DR. ZIEMER: Okay. Thank you. Okay, let's start with

 0 Henry, and then Mark and then Jim.
- good ones. I think the decision as to whether one
 would re-interview -- I think this basically sets that
 aside, because it'll depend on what the QA/QC program
 is that we need to know is where -- is a second set of
 ears sitting in and listening interacting then to
 improve the interviews as they go forward, if the set
 of ears is saying we're going to listen on every
 interview, but each interview only has 20 seconds worth
 of listening in, then you know, we sort of need to know
 is the person going to listen to the whole interview
 and then, you know, comment back to the interviewers

'cause they're all learning as they go anyway. I think part of the QA/QC how -- how it's designed would relieve a great deal of concern about the interview. And I think that's kind of why I support this language that I think what we're basically saying here is that's an option, but we aren't -- we don't need to really think about that till we see, gee, if you look at all these interviews, if there doesn't -- if they don't seem to be generating anything or everybody's in agreement, then I don't think we'd move forward. really think we need to have this for a kind of an audit trail so that when our people come in they can look at this and say yes, this one was -- somebody did listen in and so when we look at it, there was agreement between the person who listened in with the interviewer as to how the information was recorded and what was heard. So you know, our earlier discussion about should our person sit in and listen and take notes, here their internal auditor is doing that and it's a question of then seeing how is that used or how extensive is that -- that listening in and what is the feedback loop for it. So I -- I think what's here is -

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- is a good first step and it'll help us down the line when we look at inter-- an interview and say gee, you know, there doesn't seem to be much here. I wonder -or something like that. We would have that answer in whatever the audit program was, so that's why I think it's very helpful and I -- I don't -- you know, I'm --I'm assuming or I'm hoping that whatever is designed and shared with us, we would view that, after the fact, as being sufficient for us not to be concerned about the interview process. But that -- that's a -- yet to be determined till we get into actually looking at the individual cases or our contractor starts generating that to say gee, here's some improvements we might want to recommend.

5 DR. ZIEMER: Thank you. Mark?

6 MR. GRIFFON: Yeah, I guess the first thing that strikes me in this -- in these recommendations is that it -- it's really recommending internal audit as opposed to a Board audit. And Tony is right that the tasks as we laid them out for the subcontractor right now do include a review of the procedures and the interview form. And I think that'll be telling -- once we

initiate that, we may have some good input and sense of that. 3 I guess I expected that these audit recommendations and the listening in, as Henry described it, might be a Board function. I'm sorry -- and -- and then the other -you know, I would also -- I think that last language, the re-interviewing, I don't know if the notion of requiring taping -- I know we had some early discussions about that and there are some complications about that. I don't know if the working group discussed that any further. You know, pending the outcome of the initial review of the procedures and the interview form and maybe of this internal audit, you know, we may -- we may want to go to -- and I know there -- there's hurdles to get over for that, but we may want to go to a function where we tape some of these interviews and then we can audit them in that fashion instead of re-interviewing, necessarily --'cause I know there's certainly pitfalls with the reinterviewing process, but -- but - but I guess -- one think I would ask the working group is, you know, this 2 being an internal audit by NIOSH, did -- did you have

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discussions about the Board doing this -- these steps
      (a) through (f) or whatever?
3 DR. MELIUS: Yeah, let me -- 'cause --
4 MR. GRIFFON: Or maybe I just need clarification, I don't
      know.
6 DR. MELIUS: Yeah.
7 MR. GRIFFON:
              Yeah.
8 DR. MELIUS: I think if you look at this -- I may have
      mischaracterized it. One and two is the NIOSH program.
       If you go to the last -- is what NIOSH will do is a
      QA/QC program to improve -- be improving the interview.
       If you go to the last paragraph, it's what the -- what
      we're doing, the Board is doing and doing that, and
      that was really another working group and the Board
      that is laid out the parameters for that and I think
      we've talked about it at other -- other meetings.
7 And then I think we're making -- maybe the wording isn't as
      complete as it could be, but we're making a statement
      that, you know, based on this imple-- implementation of
      these two things, the NIOSH QA/QC program -- it's
      called that -- our individual dose reconstruction
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      reviews that involve evaluating the interview record,
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therefore at this time we're not recommending that there be any further -- more intrusive way of -potentially more intrusive way of reviewing the interviews, whether it be listening in or reinterviewing or reviewing a recording of the interview or -- or how -- whatever that may be 'cause they all raised a number of -- number of issues, you know, beyond what we've talked about here, so -- a big headache for Larry to deal with and some of these could be, anyway. So I think that therefore we're -- you know, this is our recommendation at this point in time, and I think we have to leave it open and see what happens down the road and see what the results of these are.

5 DR. ZIEMER: Larry, you had a comment?

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6 MR. ELLIOTT: Yes, if I might. You've heard me speak about this before, and in the spirit of being helpful and not in the spirit of belligerency or -- or unhelpfulness, these set of recommendations are appropriate, we feel, for an understanding of what it is the Board would like to audit on this piece of the process. Many of these are already in place or being developed. Yes, we want

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to be very clear that we don't have all of these fully
      developed and fully functional, but that's the
      direction that we are going. You've heard me say from
3
      the very start that we think the Board's audit should
      evaluate the interview process and how they contribute
      to dose reconstructions.
7 Now, just for your benefit, you've also heard me say that
      re-interviewing claimants is off the table. And it's
8
      not this Board's decision at the end that's going to
      make that -- that'll carry that day. The Department
      will weigh in on whether or not this actually happens.
       If you so choose to leave the door open, that's one
      thing. But if you choose to ask for a re-evaluation
3
      and re-interview of claimants, the Department will have
      to weigh in on that. So I just offer that as helpful
      perspective, not as a belligerent perspective.
      you to understand, at the end of the day the Department
      will have to decide the value of that particular
      component if you choose to re-interview claimants.
O DR. ZIEMER:
              Thank you. Okay. Let's go right down the
      line. Leon?
2 MR. OWENS: I'd just like to say that I agree with the
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recommendations by the working group. I think they've done a very good job. 3 In regard to the re-interview process, I appreciate Larry's comments, but I do think that we need to at all times consider the credibility of the program and by making that consideration with the claimants, particularly the elderly, that are not as well-versed in this process as we might be, I think there would be value-added in a re-interview from the standpoint of quality assurance. O DR. ZIEMER: Thank you. Roy? 1 DR. DEHART: We have a proposal here, and I assume that the rationale for that is that we're not sure that the current interview system is working effectively. I would ask the question, having done 20,000 interviews as was presented today, are we aware of any significant problems with the interview process? 7 MR. ELLIOTT: I would answer that question this way, that no, we are not aware of any problems in our interview process. ORAU does have the manager of that particular task and other delegated folks in that part of the program listen in -- and it's not just for 20 seconds; it's for the whole interview that -- they listen to the

whole interview and feedback is provided. 2 I will also say this, that there have been a number of cases that I have approved to go over to DOL where I have seen how the interview has been captured and utilized in the dose reconstruction, and how it's reflected in the dose reconstruction report. So I'm fairly comfortable and confident in saying to you today that this interview process is a contributing factor to dose reconstruction, and we have not identified any major problems with it. But we're watching it very closely. 1 DR. DEHART: I gather from what you had said previously that you're in the process of developing just what is being suggested here, basically -- a way of going through and documenting that the -- basically (a) through (f) -some modifications will be occurring. 6 MR. ELLIOTT: Several of these are in place. They're perhaps not at a state of readiness that we are happy with. Some -- a couple of these are not in place, but they're -- we've had similar ideas and we intend to put them into place -- (f) for instance is one -- you know, we don't have in place right now, I don't believe, 2 necessarily, but we do agree it needs to be put to --

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put in place. And the traveling document, hard copy, would go along with that, so -- appreciate those thoughts and those comments.

4 DR. ZIEMER: And if I might add, again, I just observed the work of the subcommittee (sic), I don't think there was an assumption that there was something wrong currently The real issue is how do with the interview process. we carry out our responsibility of evaluating it. And I certainly became aware as we got the materials from Dr. Toohey on -- on what they do and how they do that -- for example, they do have a management tool where they listen in to the interviews. Great, how do we critique that? If the -- is that available for us to look at so we can say what -- what was the evaluation of the listener of that interview. So these things simply reflect the kinds of records that could be audited where we could make a judgment. Yeah, the interview was properly reflected -- and all the things that Larry just described. We want to be able to -- to confirm those kinds of things, how the interviews are used in the dose reconstruction, if indeed they are; how they've contributed to it. So it's a matter of

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simply being able to document what has been talked
      about here.
3 We'll go on to Tony, and then we'll circle back again.
4 DR. ANDRADE: Okay. Perhaps -- perhaps my comments have
      been overblown a little bit, or perhaps I overblew them
      a little bit. If the process that we're suggesting
      remains internal and a re-interview is -- is -- let's
      say it's decided by one of the reviewers here in the
      list that runs from (a) to (d), determines that more
      information is necessary -- like an internal decision
      to re-interview is appropriate before the case is
      closed.
3 However, that's just -- I'd just make sure that everybody
      knows where I'm coming from. Once the Board has
      decided to take a look at these things, I understood
      that we were going to be looking at cases that were
      closed. Those are the cases that I'm concerned about.
       If they are closed, if there has been a decision that
      was not positive, then I don't want to see our
      recommendations used to try to provide an avenue to
      redress the decision. That's where I'm coming from.
2 If we're talking about the internal processes that have been
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described, then I'm fine, and this language is
      perfectly fine.
3 DR. ZIEMER: They can re-interview now.
4 DR. ANDRADE: Go ahead?
5 DR. ZIEMER:
              They can re-interview currently if they need
      more information. That's not --
7 DR. ANDRADE: As many times as they need.
8 DR. ZIEMER: -- the issue that's being -- that's not the
      issue, I don't think. I think the issue was exactly
      what you described; you want to re-interview a closed
      case.
2 DR. ANDRADE: Right.
3 DR. ZIEMER: That was the -- all right, let's go back here -
      - is it Mark next? Then Jim.
5 MR. GRIFFON: Yeah, just a couple of things. I mean I -- I
      think -- you know, one thing that I've heard in
      previous meetings from public comment is that there is
      a concern with the interview and the information that's
      being collected, so I don't know that we don't have any
      concerns over it. We've heard that expressed at
      several meetings. Maybe those things have been
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      corrected. I don't -- you know, that was a while --
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some of these were a while ago, but you know, to say
      that we haven't had any concerns over this, I think is
      not -- is not true. I think we do have some concerns
      over that.
5 The second thing I was going to ask is if the Department's
      policy is that re-interviewing is off the table, what -
      - what's the policy on this -- we -- we did bring up
      the idea of taping and creating a transcript of the
      interviews upon the consent of the claimant, obviously.
       Is that off the table? Can that be -- is that
      something that the -- the Department would consider?
      know it -- there's hurdles involved, but --
3 MR. ELLIOTT: We have considered that, and we have
      articulated the problems associated with that numerous
      times.
              And at this juncture, it's -- it's not a viable
      recourse.
7 DR. ZIEMER:
              Jim?
8 DR. MELIUS:
              Yeah, a few comments -- address some of -- some
      of these points. First, as Mark said, I mean the
      interview is a very public part of the program, and we
      are much more likely to have people concerned about
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      their interview than to ask questions about IMBUS (sic)
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or how IMBUS is calculated or -- or whatever, what assumptions were used or these other technical information. So it's always going to be very visible, and so I think in -- and people may sort of attribute more importance to it than is appropriate in their individual case or something, and particularly given the time frame involved, the survivor issue and so And so I think we have to have a credible forth. process in place. NIOSH has to have a credible program to review it and then continue to -- continue improvement issue, and then the Board has to have a credible process for -- for reviewing it, so -- read there -- and again, in going through it, we didn't see any particular problems. However, what we did notice was that there were a number of places that NIOSH had -- steps NIOSH had in place that reviewed the interview. They were listening in, there was an initial review, there's a later review. I think the question of are we going to find problem -- potentially find problems with the interview -- probably going to come up -- the issue with -- at the point of the individual dose reconstruction where someone's looking in a lot of

detail at all the information on the case and would notice discrepancies, potential problems. Those may very well be dealt with by, you know, a quick check of the record or a quick call back to the person for clarification or something, and that's fine. And all we're asking for here is really that that be reported in some way so we -- so we have a record of it and so forth. So I think that's -- go forward. 9 I think -- yeah, we understand the Department's going to be resistant to re-interviewing and so forth. At the same time, we have an obligation, you know, that Congress gave us to review the dose construction -reconstruction program, and we have to be able to say that we're -- that we as a Board are doing that properly. And if -- and that means we have to be able to say something about the interview program. And we don't want to be put in the position where we're having to say that we could not carry out our assigned mission because we weren't given the capability or the access or tools necessary to -- to review a major part of this program. I think what we've laid out here may get us 2 there, so I think that's --

1 DR. ZIEMER: I think that's a good point, Jim. I think the subcommittee (sic), at least from what I heard, felt like if there was in place a good quality assurance program and that that could be audited, in fact we would be able to reach the level of confidence that we're talking about -- a good possibility of reaching that without having to do a re-interview. would be key items that would help us get there, and that was the thrust of it, I believe. 0 Okay. Roy again? 1 DR. DEHART: 2 DR. ZIEMER: No. Okay. Gen, Gen Roessler. 3 DR. ROESSLER: I'm totally in support of the proposal here and the motion down through number two, but I'm not comfortable with even bringing in the wording about reinterviewing, for the same reasons that Tony has stated. Even though it says we're not going to do it at this time, by bringing in the wording, it leaves it open. And I just think that's inappropriate. I would totally support this if we went with Tony's friendly motion or whatever it was, but it --2 DR. ZIEMER: I don't know if Tony actually made the motion.

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You're certainly free to make the motion if you wish -
3 DR. ROESSLER: I guess what I'm saying is I'm totally for
      this if we can take out those sentences that include
      re-interviewing, even though it says we don't recommend
      it at this time. Just bringing the wording in leaves
      it open. It leaves it sounding like this is a
      consideration, and I -- well, maybe if it is later on,
      it could be brought up later on, but I -- I don't think
      the wording needs to be in it at this time.
1 DR. ZIEMER:
             And again I'll point out, I don't know if
      you're making the motion yet or not, whether it's there
      or not doesn't preclude the Board taking some other
      action at a later date in any event, whether or not you
      wanted to.
6 DR. ROESSLER: But then I --
7 DR. ZIEMER: Are you making a motion to amend by deleting
      the last two sentences?
                I could do that. Yeah, I -- I think the
9 DR. ROESSLER:
      point is, whether it's there or not, if we can do it
      later on, let's just leave it for later on. Let's not
      even bring in the thoughts at this time. I don't see
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that it adds anything, and I think it detracts from --
2 DR. ZIEMER: So are you speaking in favor of the motion that
      you haven't yet made, or...
4 DR. ROESSLER:
               I guess I was hoping Tony would make the
      amendment to the motion. I think he had the wording.
6 DR. ANDRADE: I'll let Mike --
7 DR. ZIEMER: Okay, we have a comment first from Mike.
8 MR. GIBSON: Just on the issue of the re-interview, you
      know, I don't think we're trying to say that, you know,
      NIOSH isn't doing the right thing, asking the right
      questions as they know them. But there could be some
      things that come up during a site profile that would --
      could possibly reflect back on the claimant and they
      would need to re-interview them, too, that wasn't
      necessarily known to NIOSH at the time.
6 DR. ZIEMER: Any further comments? Are there -- does anyone
      wish to amend this document before we vote on it?
                                                         Ιf
      there are no... There appears that -- okay, Tony.
      not trying to urge you to do it. You can either...
O DR. ANDRADE: I really don't want to have another meeting to
      discuss all of the intricacies here. Mike brings up a
      very good point, and there are other concerns on the
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table. But I think we've had our Federal official
      advise us about the likelihood of us ever doing
      retrospective interview after a final decision has been
3
      made, and I -- based on that, I think -- not I think, I
      will make the motion that the only change really --
      this is a -- this is a very well-done draft here, Jim,
      that the only change that I would submit for the
      Board's consideration in a motion to move on this is
      that we delete the last two sentences of the document
      and go forth with the rest as a recommendation to
      NIOSH.
2 DR. ZIEMER: Okay. This is a motion to amend the document
      by deleting the last two sentences. Is it --
4 DR. ROESSLER: I second it.
5 DR. ZIEMER:
              Seconded. Now is there discussion on the
      motion to amend?
7 DR. MELIUS: Yeah.
8 DR. ZIEMER: Okay, here and then there.
9 DR. MELIUS: (Off microphone) Yeah, a couple of points.
0 MR. ELLIOTT: Use your mike, please, Jim.
1 DR. MELIUS: Sorry, I didn't realize that Gen had borrowed
      it here.
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1 Two things. With all due respect to our Federal official, we are -- our charge is in some ways separate from them and I -- I hate to have us be doing an amendment in reference to having Larry tell us we shouldn't be -what we can and cannot do to review the program we're supposed to be reviewing. I think that raises some issues about our -- our charge. I think -- appreciate what he's telling us factually and -- and so forth and I don't think that's his motive, but I think --(Inaudible) by that.

1 Number two, my understanding of the charge to our working group was to deal with this issue, and in some sense the reference to re-interviewing is because of the way Paul gave us the charge and the discussions we had in order to carry this out. And it was specifically not to design how the program could be reviewed or develop what's the best review of the interview process, but rather were there things that we could do that would be sufficient, short of re-interviewing or some other, more intrusive process that -- to do. And I think -- I think we need to have that reference in there. think it commits us one way or the other and if there's

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wording that would, you know, add -- you know, re-
      interviewing or other methods of evaluation, that's
      fine. But I think we need that reference -- reference
3
      in there. You know, I -- quite frankly, I think the
      committee's split on this issue and -- and it's a hard
      one to deal with conceptually. I think the way we're
      taking -- again, aside from those two sentences, the
      way we're taking is a way of trying to develop a
      compromise that everybody can live with and -- and then
      when we get, you know, down the road, whatever it will
      be a year or two years when we have this information,
      we'll be able to make a more informed recommendation
      one way or the other. And maybe our differences will
3
      be less at that point in time, but I certainly feel
      that that's -- this should be -- reference should be
      kept in there.
7 DR. ZIEMER:
              Okay. So you're speaking against the motion.
      Okay.
9 MR. ELLIOTT: Point of clarification.
O DR. ZIEMER: Point of clarification.
1 MR. ELLIOTT: Point of clarification, the charge to the
      working group was specific in evaluating options or
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identifying options to evaluate the interview process.
2 DR. MELIUS: That's not the way that Paul gave me the
      charge.
4 DR. ZIEMER: Well, I don't recall the exact wording.
      tell you that the Chair's objective is to try to find a
      way to audit this without doing interviews, but -- but
8 MR. ELLIOTT: I can assure you, that's the exact charge from
      the transcript. I sent it to the working group the
      other day.
1 DR. ZIEMER: But I -- but I don't -- again, let me point
      out, it doesn't -- I'm not speaking for or against the
      motion. I would point out if the sentences are struck,
      this does not preclude anything. It simply doesn't
      address it right now.
6 Henry?
7 DR. ANDERSON: I guess one -- one argument I would see for
      leaving it in is it provides some institutional memory
      that in four years every Board member here could rotate
      off. In a year we may have --
1 DR. ZIEMER: Where did you get the four?
2 DR. ANDERSON: Well, the assignments is -- the assignments
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are one to four years. I rotate off in four months,
      and there's two others -- we haven't -- it hasn't been
      discussed here -- are slated to potentially rotate off
      or at least would have to be renominated, others would
      be -- potentially could come on, so one of the benefits
      I see of having this here is we've -- we've spent, I
      would say, almost an inordinate amount of time
      discussing it, but it isn't reflected anywhere in any
      of -- a recommendation or things we've made, so yes,
      it's there kind of lost and forever of the transcripts.
       Somebody would have to go back if, when the review
      comes up four years from now, and somebody says well,
      let's go back and see what our recommendations -- were
      they met, this would just remind you well, here's an
      option that was discussed. So that -- I think it's
      helpful to have this. It doesn't make a commitment,
      but as you say, you could go back, but somebody may not
      even remember about --
9 DR. ZIEMER: Let me point out, Henry, where this is going to
      appear and that is in the transcripts.
1 DR. ANDERSON: Well, but -- but if you --
2 DR. ZIEMER: The transcripts are there already. That's the
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institutional memory. This is -- this is no better
      than the transcripts in which it will appear anyway,
      one way or the other. The issue has appeared over and
      over in the transcripts. That's all I'm saying.
5 DR. ANDERSON: Yeah, well, I -- I -- I --
6 DR. ZIEMER: But -- no, I do point --
7 DR. ANDERSON: I'm just saying --
8 DR. ZIEMER: I understand --
9 DR. ANDERSON: -- on every other board I've been on, what
      the agency pays attention to and subsequent board
      members is what are your action items and -- and we've
      tended not to maintain a list of action items.
      have is resolutions and recommendations that we've
      made, and one could go back and say --
5 DR. ZIEMER: Gotcha.
6 DR. ANDERSON: -- as a new member, you'd want to -- you'd --
7 DR. ZIEMER: A specific action item.
8 DR. ANDERSON: You'd want to know what are the specific
      issues. That -- you know, I just raise that as one
      reason for having it here.
1 DR. ZIEMER:
             Okay.
2 DR. ANDERSON: It doesn't make a commitment to do it, but to
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have no mention of it --
2 DR. ZIEMER: Okay.
3 DR. ANDERSON: -- means somebody has to think of it later
      down the --
5 DR. ZIEMER: You speak for the motion then. Anyone speaking
      against the motion? Okay.
7 MR. ESPINOSA: Against the motion?
8 DR. ZIEMER: Well, I'm going to alternate. Against the
      motion?
0 MR. ESPINOSA: Yeah, I'm against the motion. I agree with
      Henry. I believe that it should be in there.
      doesn't matter whether it's left out or put in, either
      way it does give us something to reference back on, as
      Henry's saying. So therefore I speak against the
      motion.
6 DR. ZIEMER: Okay. Anyone for the motion?
7 DR. DEHART: For the motion?
8 DR. ZIEMER: For the motion.
9 DR. DEHART: I speak on behalf of the motion, and I'll
      explain very quickly why. The title is the
      recommendations of, and the next to the last sentence
      begins (reading) At this time, the working group is not
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recommending... 2 We're countering what we're saying in the document. I speak for the motion to delete. 4 DR. ZIEMER: Okay. Mark? 5 MR. GRIFFON: Yeah, I'm speaking against the motion. -- and one of the reason-- I mean I've already spoke to some of my concerns about it, I reflect back on a veteran's program review that John Till did and one of the findings that they had was how important the -they weren't even interviews, really, they were -- they were I guess written documents provided by some of the claimants in those cases, and the fact that the dose reconstructors may have underestimated doses in many cases because they didn't incorporate adequately the information that was identified in some of those interviews. So I think -- you know, this is such an important -- I think it's an important component. Ι think how it's designed is critical to how much information you're going to elicit from these people and how -- how useful it can be. I think it can be very important, though, and I -- I don't want to close 2 the door. In fact, I was hoping that the working group

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would come out with a recommendation on how to -- to
      re-interview or tape or -- or to actually have the
      Board do an audit. I think it's a little soft in that
      regard, but at least I want the last clause in that
      allows us -- reminds us that the door is open to re-
      examine this issue upon findings of the other steps in
      -- in section one of this document.
8 DR. ZIEMER: Just for clarification, Mark, wasn't Till's
      concern that material elicited in the interviews wasn't
      in fact utilized, as opposed to the issue of not having
      adequate interviews? See, I --
2 MR. GRIFFON: Well, yeah -- yeah, they didn't have
      interviews, but the -- yeah. I mean but the -- they
      said that they weren't -- that they weren't really used
      by the dose reconstructors adequately and they didn't
      pay attention to them or --
7 DR. ZIEMER: But the issue that we have is sort of a
      separate issue.
9 MR. GRIFFON: It is sort of a separate issue, but it just
      demonstrates the importance of the tool, is what I'm
      saying.
2 DR. ZIEMER: The interview is definitely important.
                                                      That's
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why we're talking about reviewing it. Okay. Wanda has
      a comment.
3 MS. MUNN: I need one point of clarification. We are
      talking about closed claims. Correct?
5 DR. ZIEMER: Closed claims. That's correct.
6 MS. MUNN: If we are talking about closed claims, then the
      consideration of re-interviews puts this Board in the
      position of being viewed by the public as a quasi-
      appellate group.
0 MR. PRESLEY: That's exactly right.
1 DR. ANDRADE:
               Exactly.
2 MS. MUNN: I am not prepared to serve as an appellate. I
      don't believe that was the charter of this group and I
      won't go there.
5 DR. ZIEMER: Thank you. Further comments? Roy, you have
      another comment? Robert?
7 MR. PRESLEY: Well, early on we were in Cincinnati and some
      of us witnessed some of the interviews. And I'm going
      to be honest with you. I'm -- I'm accepting to what
      they're doing. I think if we go to this, we're getting
      ourself in trouble.
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2 DR. ZIEMER: So you speak for the motion?

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1 MR. PRESLEY: For the motion.
2 DR. ZIEMER: Any other comments, for or against the motion?
       Are you ready to vote?
4 Okay, those in favor of the motion, raise your right hand.
                    (Affirmative responses)
6 DR. ZIEMER: One, two, three, four, five for the motion.
7 Those opposed?
                     (Negative responses)
9 DR. ZIEMER: One, two, three, four, five, six opposed.
      Chair votes for the motion. We're six and six.
      Chair rules that the motion does not pass. It requires
      a majority to pass. That's not the same as failing,
      but it doesn't pass. Which means we return to the
      original document, as written. Okay?
5 Now incidentally, under Robert's Rules, anyone can challenge
      the Chair's ruling, and the challenge has to be upheld
      by two-thirds vote, so -- so the ruling is that the
      motion fails for lack of a majority. It seems like the
      Chair ought to get two votes in these cases.
0 DR. ANDERSON: Actually I thought the Chair didn't vote
      unless it was a tie, but that's okay.
2 DR. ZIEMER: Actually under Robert's Rules, that's generally
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the case. But I think we agreed in our own rules that
      the Chair would always vote so that people knew where
      the Chair stood or didn't stand.
4 So the motion fails and we have before us the original
      document, as written. Is there any further discussion
      on the document as written?
7 Are you ready to vote on the document which -- the
      recommendations and the not recommendations of the --
      usurping Roy's characterization.
0 A comment, Tony?
1 DR. ANDRADE: Just a quick question for you, Paul. As I
      mentioned, I didn't have a chance to provide my
      editorial and lesser comments to Jim.
4 DR. ZIEMER: Can we do these as friendly amendments? Are
      they -- are they strictly editorial?
6 DR. ANDRADE: They really are.
7 DR. ZIEMER: Do you want to --
8 DR. ANDRADE: Would it be possible to do that, even if we
      adopt --
O DR. ZIEMER: Yeah, let's do it real quick.
1 DR. ANDRADE: -- the document as written?
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2 DR. ZIEMER: Just tell us what they are.

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1 DR. ANDRADE:
               Okay.
2 DR. ZIEMER: We'll see how friendly they are.
3 DR. ANDRADE: Okay, on point number 1(a), records of the
      current -- like I said, this is editorial -- management
      monitoring instead of supervisory monitoring.
6 DR. ZIEMER: Any problem with that, management monitoring?
                        (No responses)
8 DR. ZIEMER:
              So ordered.
9 DR. ANDRADE: Okay. On point (b), the second step in the
      review is not necessarily done by management, but it is
      done by another group of people that look for accuracy
      in -- both technical and editorial. And so I'd like to
      strike the word "management", and also call it the
3
      proper -- give it the proper title, and it's review of
      the -- and it's not completed interview, it's the
      summary report to...
7 DR. ZIEMER: Is this the review that -- where somebody's
      looking at it for everything from grammar to --
9 DR. ANDRADE: Yes.
O DR. ZIEMER: What is the proper terminology we want here?
      Can on staff help --
2 MR. ELLIOTT: He's got it.
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1 DR. ANDRADE: It is the summary report.
              It's the summary review, okay.
2 DR. ZIEMER:
3 MR. ELLIOTT: Summary report.
4 DR. ANDRADE: To, and then "include" instead of "including"
6 DR. ZIEMER: The summary review of the completed interviews?
7 DR. ANDRADE: Of the summary report --
8 DR. ZIEMER: Summary report.
9 DR. ANDRADE: Instead of completed interviews, to include --
      and then here's an insertion, "items that are found to
      need further clarification" and strike everything up to
      "and corrective actions".
3 DR. ZIEMER: Okay. Let me ask the working group, are you
      comfortable with this rewording as far as --
5 DR. MELIUS: That's fine, yeah.
6 DR. ZIEMER: -- it covers the intent still? Thank you.
7 So is it -- let me read what I have and see if it agrees.
      Records of the summary report of the completed
      interviews, to include --
0 DR. ANDRADE: No, no, no, no.
1 DR. ZIEMER:
              No?
2 DR. ANDRADE: These are not completed interviews.
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1 DR. ZIEMER:
              I'm sorry.
2 DR. ANDRADE: These are still in review process.
3 DR. ZIEMER: Read it again. Read it again, I --
4 DR. ANDRADE: Okay. Records of the review of the summary
      report, to include items that are found to need further
      clarification, including corrective actions. So by the
      word "items" I'm including anything small or large,
      including significant problems. Okay?
9 DR. ZIEMER:
              Okay.
O DR. ANDRADE:
               Item (c), second line there, "dose
      reconstruction is being done to include any items --
      instead of "significant problems" -- found --
3 DR. ZIEMER: That is a slight -- that is probably slightly
      more than an editorial. It changes the level of
      findings, but let me ask if the working group considers
      that a friendly amendment or do you have prob-- it
      actually requires more reporting than you would have
      suggested here, I believe. It lowers the bar a bit on
      what's reported. Working group okay? Chair of the
      working group?
1 DR. MELIUS: Yeah, that's fine. That's not...
2 DR. ZIEMER: Without objection, we'll consider that a
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friendly amendment.
2 DR. ANDRADE: Okay. To continue on with that, "to include
      any items found" -- and then again an insertion -- "to
      need further clarification" -- and then we continue
      with the phrase, "and corrective actions".
6 DR. ZIEMER: Ray, did you get all that, as well, for the
      record?
               Thank you.
8 DR. ANDRADE: The very last editorial comment is in the
      phrase that falls right underneath -- right underneath
      (f). And frankly I didn't have a chance to construct
      it, but I wanted to include the fact that we are having
      a subcontractor perform part of this review, and I just
      didn't know exactly where to fit that in there, Paul.
4 DR. ZIEMER: Notice that (a) through (f) are things that
      it's suggested be part of NIOSH's system so that we can
      review it, so I'm not sure we'd bring our subcontractor
      into the picture at this point, if I understand what
      you're saying.
9 DR. ANDRADE: Okay. Well, maybe that's appropriate to just
      leave it off.
1 DR. ZIEMER: This is just identifying sort of a body of
      kinds of records that could be reviewed, I believe.
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1 DR. ANDRADE: Okay. I'll agree to that.
2 DR. ZIEMER: Let me ask again, is the Board comfortable with
      these friendly amendments to the document?
4 MR. OWENS: (Off microphone) On (c) --
5 MR. ELLIOTT: Can you speak into the microphone?
6 MR. OWENS: Oh, I'm sorry. Tony, on point (c), in the
      previous (b) we changed completed interviews --
8 DR. ANDRADE: Right.
9 MR. OWENS: -- to summary report.
O DR. ANDRADE:
              Right.
1 MR. OWENS: And so in (c) are we going to leave completed
      interviews or are we in essence going to change that to
      summary report, also, to make it consistent?
4 DR. ANDRADE: Good catch, Leon. It should be summary
      report.
6 DR. ZIEMER: Well, let me follow up. What is it that the
      health physicist is using, the summary or the
      interview? Is it -- do you know -- can -- who can
      answer that? One of the staff. Is -- this is saying
      records of the health physicist's review of the
      completed interviews at the time dose reconstruction's
2
      being done, so what is it the health physicist is using
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at that point, is the question.
2 MR. ALLEN:
            Paul --
3 DR. ZIEMER:
              Yeah.
             Well, the completed interview we call the
4 MR. ALLEN:
      summary report of the interview.
                                        It's a summary
      because it's not a transcript. The --
7 DR. ZIEMER: So the health physicist is using --
8 MR. ALLEN:
             It's the completed --
             -- the summary -- the completed summary report.
9 DR. ZIEMER:
0 MR. ALLEN: Yes, it's the completed one. In item (b) I
      think it's -- I think the difference is it's kind of a
      draft at that point. It hasn't been reviewed.
      could possibly change for grammar, et cetera.
      that make any sense?
5 MS. MUNN: Uh-huh, yeah.
6 DR. ANDRADE: Yeah, it's a little confusing, but --
7 MR. ALLEN: Yes, it is.
              -- recall that the health physicist himself
8 DR. ANDRADE:
      can review this first report that comes down to him,
      which is still basically a summary report, and still
      ask for more information and really actually ask for a
      re-interview if necessary.
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1 MR. ALLEN: Right, but the difference at this point is (b)
      is before the claimant ever sees it. After that
      review, then it goes to the claimant, the claimant can
      make any changes he wants to. In (c) the claimant's
      already seen it, has made any changes they want to, and
      the dose reconstructionist is the one looking at it.
7 DR. ANDRADE: So if the health physicist -- if the health
      physicist has a question and say brings up a small
      technical point that needs to be clarified, and that
      may require an interview again or may need some
      clarification from other people that have listened in,
      then what is it called?
3 MR. ALLEN: At that point, if we were to call the claimant
      back again or get a letter or whatever from the
      claimant to change anything, we change it and it's an
      updated summary report, is what we call it.
7 DR. ZIEMER: But for practical purposes, it's the completed
      summary report --
9 MR. ALLEN: Completed summary --
O DR. ZIEMER: -- or completed interview report. Is that the
      same as --
2 MR. ALLEN: Yeah, there's very little -- the semantics
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aren't too --
2 DR. ZIEMER: I think we understand it then.
3 MR. OWENS: And I didn't intend to do anything --
4 MR. ALLEN: In any case we'll get the right seman-- the
      right titles --
6 DR. ZIEMER: We'd better vote before... Are you now ready
      to vote? And because of the previous vote, you
      recognize -- and I'm recommending that those who voted
      for the prior motion not vote against the document
      because of the presence of the last two sentences,
0
      recognizing that the record will indicate that there
      was a sort of split on the issue of the last two
      sentences, but the rest of the document perhaps could -
3
      - not that the Chair's trying to influence anybody, but
      we -- we don't want to throw everything out, if
      possible. We need the rest of the document, so --
7 MR. PRESLEY: Clarification?
8 DR. ZIEMER: Yeah.
9 MR. PRESLEY: Last paragraph, third line, "At this time the
      working group", should we not change the working group
      to Board?
2 DR. ZIEMER: That's a good point. I think a friendly
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amendment here needs to be made. Does the subcommittee
      (sic) agree? It now is a recommendation of the Board,
      not the subcommittee.
4 MR. ELLIOTT: And also up in the first sentence.
5 DR. MELIUS: And the first sentence should read the same,
      too.
7 MR. PRESLEY:
              Yes.
8 DR. ZIEMER:
              Thank you.
9 MR. PRESLEY: Yes, I...
O DR. ZIEMER:
              Ray, that's the first sentence in the document
      will read -- instead of working group, Advisory Board
      on Radiation and Worker Health, and then the third line
      of the last paragraph, instead of working group will
      read Advisory Board on Radiation and Worker Health.
5 Now are we ready to vote? All those who support the
      document, please say aye.
                    (Affirmative responses)
8 DR. ZIEMER:
              Those opposed, no?
                         (No responses)
O DR. ZIEMER:
             Any abstentions?
                         (No responses)
2 DR. ZIEMER:
              The motion carries. Thank you very much.
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1 MR. ELLIOTT: I'd like to thank the working group for their
      efforts on this.
3 DR. ZIEMER:
             Appreciate it. Okay, we're a little over time
      for the break. Let's go ahead and take the break.
      Break till about 3:00, if that's agreeable.
6 Let me also ask -- is Cori here? Do we have anyone signed
      up for public comment for this afternoon and how many
      individuals? Two individuals? If the individuals -- I
      want to ask if the individuals -- the public comment
      period is scheduled for 2:45. We can -- we can go
      ahead with that now or we can take the break.
      depends on whether those individuals need to leave or
      would just as soon have the break themselves right now.
4 DR. MELIUS: Let's break and...
5 DR. ZIEMER: Patricia Ehlmann and Knute Ringin -- is it
      Ringin? Patricia, are you all right if we go at 3:00
      instead of 2:45?
8 MS. EHLMANN: (Off microphone) That's fine.
9 DR. ZIEMER: All right. And Knute -- is it Knute?
0 MR. RINGIN: (Off microphone) It's Knute.
1 DR. ZIEMER: Is that -- is that okay?
2 MR. RINGIN: (Off microphone) Sure.
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1 DR. ZIEMER: Okay. Then we'll take a 15-minute break.
      Thank you.
3 (Whereupon, a recess was taken.)
                     PUBLIC COMMENT PERIOD
5 DR. ZIEMER: We're going to reconvene and proceed with the
      public comment period.
7 We're going to begin with Patricia Erlmann -- do I pronounce
      that correctly?
9 MS. EHLMANN: Ehlmann.
O DR. ZIEMER: Ehlmann.
1 MS. EHLMANN: It's German.
2 DR. ZIEMER: It's a Deutsche name. She's from Wright City,
      Missouri, which is in the St. Louis area, was not able
3
      to be with the Board when we were in St. Louis, but
      we're pleased that she's able to be here today.
      Patricia, we'll -- if you will use the podium here --
      or okay, the mike there is on, that's fine. Thank you.
                            (Pause)
9 DR. ZIEMER: Okay, thank you for your patience. Patricia,
      please proceed.
1 MS. EHLMANN: I just want to say that my name's Patricia
      Ehlmann, and my brother and I are survivor claimants
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for Everett Powers, tracking number 10141, case number 21496, file number 488092991. My dad was employed at Mallinckrodt from the 4th month of 1943 to the 10th month of 1966. He started at the St. Louis plant and was transferred to Weldon Springs around 1957 when the plant opened. He was diagnosed with multiple myeloma in 1983. His cancer was chemo-resistant, although he did have to go through a lot of chemo treatments. 9 This type of cancer attacks the bone marrow, so his bones disintegrated to the point of vertebrae fractures, which were extremely painful. During all of this treatment he also had at least two chemical peels of his head. As you can see from his picture, he was very bald, so he had a lot of skin to come off the top of his head. 6 This is a terrible ordeal to go through for skin cancers. Squamous cell cancer took part of his nose, most of one nostril and approximately one-half of his lips. the weakness of bone disintegration, he fell one morning getting out of bed -- and this was on a carpeted floor -- causing a brain hemorrhage for which 2 he had brain surgery. So now he had three holes bored

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in his head. And since this is basically a stroke, he
      had to completely go through therapy to regain speech,
      movement of his arm and his hand.
4 My dad smiled all the time, and right up to the time the
5
      morphine completely knocked him out, he was smiling.
      He died in September of 1987. My mother filed the
      first claim in 2001 and at her passing in September of
      2002, my brother and I sent in claims.
9 We're just normal people who know very little about
      government paperwork, and boy, were we in for a shock.
       After finally finding everything we could, with a lot
      of help from Paducah, we were in for another problem.
      The paperwork showed -- proved to the DOL all of the
3
      cancers that I listed above, but it only showed that he
      worked at the Weldon Spring plant. After many phone
      calls, with the help of Denise Brock, I finally had a
      conference call between myself, Ms. Brock and Jeremy
      Stanton of the DOL. He stated that their paperwork
      showed that he only worked at Destin (sic) in St.
      Louis. Now I mean this is getting crazy. I've got the
      same Department telling me he worked in two different
2
      places. We need help to resolve this. We're not the
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only ones with this problem. I have heard of other
      people where the paperwork is not showing where they
      worked. We can't get our interviews.
4 We are due an interview now because I know Destin (sic) is
      taking the interviews. He worked there long enough to
      get that interview. And it takes only common sense to
      know that if he started in '43 and he quit in '66, he
      worked at both plants because they weren't in operation
      at the same time.
O Thank you, and I'd appreciate your help.
1 DR. ZIEMER:
              Thank you. I'd like to see if any of the Board
      members have questions for Patricia.
                        (No responses)
4 DR. ZIEMER: Thank you very much. Then Knute Ringin, who
      is from Seattle, Washington. Thank you. Knute?
6 MR. RINGIN: (Off microphone) Thank you very much for
      allowing me to speak here today. With respect, I'd
      like to ask for equal time (Inaudible) that Richard
      Miller has had in front of you over the last 18
      meetings --
1 THE COURT REPORTER: Dr. Ziemer, he's not feeding.
2 DR. ZIEMER: We need to -- yeah, you may have to -- you may
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have to do your request over. 2 MR. RINGIN: Well, I'll try to keep it a little bit shorter. As I said, my name is Knute Ringin and I'm glad to be I've not bothered you at previous meetings, but I think I have enough to talk about now that it's worthwhile to take a little bit of your time. 7 I am the science advisor to the Center to Protect Workers Rights. CPWR is a non-profit research and development 8 arm of the National Building Trades department of the AFL/CIO. Since I'm going to talk about ethics a little later, I would like to make a couple of disclosures to begin with. 3 First of all, CPWR has a very large, significant and longstanding partnership with NIOSH in the area of construction safety and health. We also have a contract with OCAS to try to develop better dose and radiation monitoring estimates for construction workers. We're responsible for -- involved in medical screening programs for construction workers at Hanford, Savannah River, Oak Ridge, Portsmouth, Paducah and Amchitka. And in the course of the last six years, 2 we've probably interviewed more than 10,000 workers in

those sites. 2 We also have a contract that DOL just asked us to take on to help them establish employment verification where DOE cannot verify that a worker has been employed at a DOE site. For construction workers, that's close to 20 That's 20 percent where DOE does not have percent. employment history, let alone radiation dose monitoring history. 9 Now I'm not an expert on radiation, and I don't come here to talk to you really about radiation, radiation monitoring or radiation biology. My comments are specifically limited to construction workers, which we represent and which I know something about, and to the claimants who are construction workers. My comments may or may not be relevant for other types of claimants, but they're relevant for those workers who come as construction workers and who have some unique characteristics. 9 The first is, they're all employed intermittently at DOE facilities. The second is that they're working largely in uncharacterized or inadequately-characterized 2 environments. And the third is that they work under

uncontrolled working conditions with little or no supervision most of the time. 3 Now these workers happen to have a large stake in your program that you're reviewing. There've been many more construction workers at DOE facilities than production workers, which most people don't realize. instance, at Hanford we estimate there are 59,000 construction workers at risk for radiation exposure; 37,000 at Savannah River; more than 30,000 at Oak Ridge. And roughly half of the current claimants in your program are construction workers or their survivors, so it's not going to be an incidental issue to your deliberations. 4 We're grateful that NIOSH is finally starting to process cases. It's obviously long overdue. We're deeply concerned about that because we hear it from our members all the time. But the 1,000 cases processed so far cannot be used to est-- make any kind of determination or estimate about what this program is, should or will do in the future. So far what you've seen are the easy, the straightforward cases mostly; 2 the ones who are obviously yes or obviously no and that

have been processed pretty fast. What I'm concerned about is the 30 or 40 percent of our members where there are no valid dose data and where claimants have difficulty recalling their work history for you, and that's a large number of them. 6 Let me also say from the start that we did not agree with NIOSH's interpretation of the law and its plans for dose reconstructions, and we've had many discussions with the NIOSH management about that, with Larry and so We didn't think that this approach was going to work for our members, and so far we don't see much evidence that it is. The problems that we look at here can be traced to two fundamental flaws in the way that I think the program has been set up. 5 For claimants like ours who have problems with dose information, the original dose reconstruction rule under which the program operates is extremely lacking in specification. It's fairly specific when it comes to dealing with workers who have dose, and then it has two very general paragraphs dealing with workers who don't have dose, where they say we may either 2 extrapolate from workers similarly situated, or we may

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use some other kinds of environmental data.
      essentially all the dose reconstruction rule says. And
      it doesn't give us enough specificity to determine with
      accuracy two major effects.
5 Because we don't have defined benchmarks, it's difficult --
      maybe even impossible -- to make an objective
      determination about the completeness of the dose
      reconstruction once it's done. I don't know what we
      would compare it against for these workers.
      secondly, it ends up placing what I think is an
      unreasonable burden on the claimants to document their
      exposures and to verify the completeness of the dose
      reconstruction once it's done. These workers need
      help, and I'll get back to that.
5 And the second very big flaw in the program that you all are
      very aware of is that the administrative structure now
      is so rife with potential for conflict of interest that
      it's at this point eroded to a great extent the
      confidence in NIOSH's objectivity, particularly
      confidence among our claimants, our members who are
      claimants. And we hear it every day.
2 And it comes largely from two -- there are two effects that
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2

arise from this structure. Even though policies and procedures to prevent conflicts of interest have been developed, we're already seeing evidence that they're not adequate -- at least I think we're seeing that. And as a result of this, as I said, NIOSH has a low level of credibility among claimants that it definitely has to overcome. And I think the role of this Board is incredibly important to establish credibility for this program. That's why we have this review. And not just to review what is being done, but to review it in such a way that you re-establish credibility in the program as it goes forward.

3 I want to put this in context by using the Savannah River Site history profile as an example. My comments will reflect a meeting that we had in August on November 11 where Jim Neton and four contractors came to review the site profile document with local unions, and we appreciate that he came there on a Federal holiday, which was the only time when we could get everybody together. And everybody that was -- there were 18 local union leaders, the leaders of every union that represents workers at Savannah River. We also had a

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dozen or so workers who spanned the entire history of
the Savannah River Site, who had spent considerable
time there, to talk about their experience. And we had
three technical experts. In addition to myself we had
Jim Platner* who is a Ph.D. radiation biologist and Don
Ellisberg who's a lawyer with great knowledge about
workers comp and who was in charge of all of the
Workers Compensation programs at the Department of
Labor in the past. So we provided this kind of
expertise and we had a very good meeting with Jim Neton
and his staff.
In fact, I would say that's by far the best meeting that

fact, I would say that's by far the best meeting that
we've had with NIOSH on this program because it was a
very open give or take where they said this is what
we've done; do you think it's adequate, is it going to
work for your members and what can we do better, and we
talked about some of those things. And I'm going to
talk to you -- show you a little bit about what we told
them so that you understand why it's important to look
more carefully at these documents.

1 I appreciate the complexity of trying to characterize 50 2 years of production at Savannah River in a short

document. Anybody who's been at the Savannah River Site or Hanford or Oak Ridge or INEEL will know that these are incredibly complicated sites, and here we have a document where the text is basically 100 or so pages and a bunch of appendices attached to that that's supposed to somehow explain everything that's happened in terms of radiation to those workers who were there. That's a monumental task and a very difficult one and a very important one. 0 Yet in the end of this, we still have to ask this very basic question. The document that comes out, which ends up being a summary, a distillation of all kinds of stuff, does it end up being fair to the claimants. Is this a document fair to all of the claimants, given that there all kinds of different classes of claimants who have been workers who it's supposed to cover. 7 I don't know if it's fair to ask you all, but how many of you have read the Savannah River site profile document? Yes, I don't blame you if you haven't. It's not easy reading. It's a very dense and complicated document that at least it's taken me a long time to get through. 2 And now we have many more of these things to go

through, and I'll get back to why that's also at issue. 2 The Savannah River Site history was issued in the summer of -- during summer of 2003, but we didn't know it until actually your meeting in August, I believe, when it was announced. We were not aware that it was being developed or that it had been issued. And although we can't be completely sure of this since we don't have quite an accurate or at least consistent description of what it's going to be used for, we think this is probably a very important document, that all of these site profile documents are probably very important. Our impression is that it's been developed pursuant to the dose reconstruction rule with a purpose of, to quote the document itself, "to evaluate both internal and external dosimetry data for unmonitored and monitored workers" -- sounds like pretty much everybody, I guess -- "and to serve as a supplement to or substitute for individual monitoring data". a very large charge to this document, not very specific, and we'll only learn how it'll be used as it gets implemented.

2 When we reviewed this document we became very concerned for

five basic reasons. There really isn't a methodology in this document. If we were to try to reconstruct this document, we would not be able to do so accurately, I don't think. I don't think we'd come up with the same result. And if there's any basic issue in science, it has to be replicability. In order to get to replicability, you've got to have a method and you've got to have documentation that you can follow, and that documentation and method is not sufficient to do that. 1 The report doesn't describe the methods or documentation adequately. And as I said, as a result of this, it's not possible -- I don't think -- to replicate it. But 3 we know it was done by an ORAU contractor team that talked and spent lots of hours with site personnel at Savannah River and used largely internal documents from the Savannah River Site, without specifying necessarily why, whom or when they talked to these people. don't have the documentation about exactly what was done. And that doesn't give us a lot of comfort in the process.

2 For instance, the contractor has developed a methodology for

extrapolating maximum dose from source terms, and that's used to estimate exposure from the airborne and resuspended I guess exposures. This methodology is listed in the bibliography, but it's only listed as an unpublished document that was prepared by the contractor for the purpose of doing this project. until one has that document and reviews it, you can't understand what's been done in the report itself. becomes a little bit of a house of cards. O There seems to be some significant omissions, as near as we can tell. Now we based our review on this on a very large site history inventory that we've developed at the University of Cincinnati, and Dr. Eula Bingham has that, as well as approximately 2,000 interviews with workers -- construction workers who have been at Savannah River, and here is what we found that's significant. 8 We found that we have 83 significant site history documents that are not in -- referenced in this document and I don't think have been used. Secondly, we looked at one specific area, which is the area that's common to all 2 of the reactors used at Savannah River and there NIOSH

lists that there are 32 core radionuclides, yet we have identified at least ten additional radionuclides. if you're looking for source terms to extrapolate from, then I think it's important to include all of those. Maybe we missed something, I don't know, but that's what we think is there. 7 There's virtually no description of deficiencies in radiation monitoring programs. If you read this document, you would think that radiation monitoring at Savannah River had been just about perfect from the That's not what our workers, our members, tell us in the interviews that we have done and that we can There was extensive testimony also on document. problems in monitoring practices when DOE held town meetings in Aiken in 1999 in development of this program, and that's not at all referenced, or I don't think has been reviewed in the process of developing this document. Also the tiger teams that did their investigations of Savannah River in 1990 documented very extensive deficiencies in monitoring practices, and they're not listed.

2 There is no consideration of radiation incidents or

accidents, as near as I could tell. We've identified approximately 76 accidents over the history of the site that we do not see referenced in this report, and that I believe are important if you're going to look at exposures. 6 Throughout the document there is no apparent awareness that construction workers may have very different exposure patterns from production workers. For instance, in applying the model for extrapolation from source term to resuspended radiation, there is no consideration of something as simple as digging in the dirt. it's assumed that that dirt's not going to be disturbed I don't know where -- how these resuspended very much. -- this resuspended radiation is going to become airborne again, but there certainly isn't taken into account that it might be a source of exposure for people who dig in it, which just about all construction workers do from time to time on these sites. 9 The third concern that we have is one that we take very seriously, and I'm sure you do, and I don't like it. There is a guy named Dr. Eugene Rollins who's listed on the ORAU web site as the key -- as a key person working

on this report. He apparently also developed the model to estimate maximum dose from source terms, an important internal report that is referenced extensively in the document itself. According to his conflict of interest statement that's also listed on the ORAU site, it says that he previously worked at Savannah River, including six years in human health risk assessment and one year as a shift supervisor in health physics and radiation monitoring. Now I'm not saying that that leads to a conflict of interest by itself, but I believe according to the policies on conflict of interest, people who have worked on sites should not be working in these documents, and that's at least what the folks who came down to Savannah River told us, that was their intent. So that's something that has to be looked at very carefully because it gets at the heart of this credibility problem. 8 The fourth concern we have is a minor one, but it's technically important, and that is that we think there's a conflict with the dose reconstruction rule in this document. There's a small throw-away thing in 2 there somewhere about a curious minor adjustment to

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dose for claimants who eat wild game taken in the Savannah River area. Now I don't know what the implications are of this adjustment, but I do know this: According to the dose reconstruction rule, it allows for only one adjustment to dose -- we've talked about this many times -- and that's for smoking in lung cancer. NIOSH agreed from the start that it would not make any of the adjustments, for instance, that NCI suggested for diet and that has been used in some of the other radiation reconstruction programs. believe that's included -- at least it's an intent of the dose reconstruction rule that that should not be included, yet there is a conflict here that I think is a technicality, but that could be important. That is that if a document is to be developed pursuant to the dose reconstruction rule, then it should follow the rule itself.

8 Finally, there is no independent review of these documents, as far as I know. Before this document was put on the web and issued for use, there was no review of the underlying methodology, and in this case the unpublished source terms extrapolation method, and

there was no independent review of the document itself. 2 Now follow our meeting with the folks in August with NIOSH, we agreed to make available to NIOSH all the documentation that we have, and we would have done that a long time ago had they asked us. I don't know why they didn't come to us when they were developing the report, and it seems that this is particularly serious given that they obviously didn't hesitate to meet with the DOE site personnel. The one-sidedness of this kind of contact does nothing to dispel the sense held broadly that NIOSH is not particularly aboveboard in its work. 3 Now we have good reason at Savannah River to be concerned when you just deal with stuff that the site administration gives you. When we started our medical monitoring program at Savannah River, the site strongly objected to us testing people for beryllium exposure. They said there had absolutely never been any beryllium used at that site, and they said the same thing to NIOSH. After having tested some 2,000 people or so down there, we have about one and a half percent of the 2 workers who have been there testing positive on DOE's

recommended beryllium test. And upon those initial findings, the site administration finally 'fessed up and said yeah, there may have been a little beryllium here after all. So it's not comforting to say that it's enough to go to the site administration and ask what kind of documentation do you have; just give us your stuff or tell us what you've been doing and we'll make use of that. I'm not very comfortable with it. 9 Finally, let me also make a point about our claimants that I've made many times before. After the site -- NIOSH placed the site profile document on its web site, it invited comment on it, which one can discover by reading the web site. Apart from this being after the horse left the barn, so to speak -- the document had already been issued -- it clearly places the burden on the claimants to show that there are deficiencies in this document. And that points to me to what seems to be a very unfair balancing act. On the one hand we have the site profile reports. These are very complex documents, presumably with far-reaching significance, presented very much or pretty much as final by NIOSH 2 when it puts it on the web site. NIOSH has major in-

house expertise, vast resources through its subcontracts and so on, to put into the preparation of this thing. On the other side are the claimants. these are, by definition, either workers with cancer or their survivors. They're mostly old and frail, and they have no support. And they're expected to challenge these documents and to say hey, we don't think there's -- there are problems with this dose recon-- or with this site profile that you've done. That's an unreasonable burden to place on them, because if they don't challenge it right now, NIOSH will do nothing to change these documents. 3 This is not the only burden (sic) where NIOSH places an undue burden on claimants. You talked about the interviews earlier today, and I can safely say that our members tell us that they have trouble following the interviews that are done by phone, and many of the NIOSH interviewers have said the same thing. unreasonable to expect these old construction workers to recall a lifetime of information about radiation exposure. But much more difficult are the interviews 2 with survivors, and they're close to half of the

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claimants here. I believe that the NIOSH interviewers
      agree that they get very little or nothing out of most
      of the interviews that they do with survivors.
4 Based on this, I would ask this Board to do three things, or
      consider doing three things. One is to recommend NIOSH
      that it issues a replicable method for the preparation
      of the site profiles, and that this includes validation
      of the information that it receives from the sites,
      validation both in terms of the accuracy of what it
      receives and whether it's complete.
1 Secondly, recommend or require that independent review of
      these site profiles be conducted before they're issued
      -- and I was going to call this peer review, but I'm
      not comfortable with that term in this context because
      peer review might simply mean a health physics review,
      and I think there's something much more important.
      It's not just understanding health physics, but it's
      also understanding how it applies to workers as they do
      their work on these sites.
0 And thirdly, I would encourage NIOSH to provide claimants
      who want it or need it with much more independent
      assistance with their interactions with NIOSH.
                                                       NIOSH
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wants to be claimant-friendly. The site profile document uses this word extensively. But it fails to provide the weakest of claimants what they most need, which is an independent, knowledgeable and forceful advocate in the process of doing all of this work, and it's certainly something that we would like to work on. 7 And in addition, if I may be so bold, once you hold these meetings in the future, I suggest that you make two 8 minor changes, particularly when you're planning to hold your meetings in the tri-cities and in Augusta. First send out a notice to all the claimants who live in that general vicinity, say within a radius of 50 or 80 miles or something like that, informing them that 3 you're going to hold this meeting. We will certainly be glad to notify everybody in our programs about it. And in addition, hold an evening session for public comment, maybe on the first day of the meeting, because most people will not come to a day meeting for two very significant reasons. Either they're very old or frail claimants -- whether they're the workers or their survivors -- and they usually need help to come to 2 meetings and they rely on their kids or something like

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that, and those kids are working. Or they're
      survivors, who usually are also working, they're the
      children and is the one who have jobs themselves and
3
      can't get away during the day, so maybe a two-hour
      evening session would be, I think, very enlightening to
      you. At least if you come to Hanford. I'm sure you'll
      get an earful.
8 And with that, I thank you for your time and your attention.
9 DR. ZIEMER: Thank you, Knute. Let me ask if any of the
      Board members have questions they may wish to address
      to you before you leave the podium. Any points of
      clarification? Henry?
3 DR. ANDERSON: Yeah, what -- what kind of follow-up have you
      had from NIOSH after your meeting in Augusta --
5 MR. RINGIN: Well --
6 DR. ANDERSON: -- (Inaudible) made your other
      recommendations as --
8 DR. ANDERSON: -- Larry called us about a week afterwards
      and we had a conversation where he asked us to submit
      all of our information and documentation, which we were
      planning to do anyway, and which we've started doing
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      earlier than we had planned to do it so that we're
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providing them information initially on Hanford,
      Savannah River, Amchitka in Alaska and the Nevada Test
      Site and Oak Ridge, and that includes all of the
      information that we have in our site history
      repository, as well as all of the results that we have
      from the -- from the interviews with workers.
7 I can also say to you that doing interviews with workers,
      particularly construction workers, is very difficult.
8
      It took us a long time to learn how to do it, even
      though we're kind of specialists on construction
      workers, that unless you put it in terms that they're
      used to, occupational terms -- you know, the kind of
      tasks that they do and that kind of stuff -- we don't
      typically get very far with it. If you put it in terms
      that they understand -- we use retired construction
      workers to do those interviews for that reason -- then
      we get a lot of information.
              Thank you. Other -- here's another question,
8 DR. ZIEMER:
      Rich Espinosa.
0 MR. ESPINOSA: You had mentioned 20 percent of the
      construction workers that are claimants haven't been
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able to verify employment. How is...

1 MR. RINGIN: When the Labor Department gets a claim, it sends it -- along with a form called the EE-5 form -to the local Department of Energy site where the worker has been working. Our in our -- for our workers, you know, there are many different sites. The DOE facility is then supposed to establish whether or not they have records indicating that this person was an employee on their site. For construction workers, in about 20 percent of the claims that DOE site that they send that record to is unable to verify in its documentation -one of the problems is that a lot of our members have been employed by subcontractors and sub-sub tiers of contractors, and it becomes -- having records. Now 3 there should be records somewhere. It's -- you know, you're required to submit certified payroll records every week or month on all of these workers, so somewhere it should be, but they are unable to come up with it in a timely fashion. 9 DR. ZIEMER: Thank you. Other questions? O MR. ESPINOSA: I got... 1 DR. ZIEMER: Oh, a follow-up. Okay.

2 MR. ESPINOSA: One of the things that concerns me,

especially when it comes to the SEC, is the co-worker data, such as if they didn't have enough data for me, they're going to use a co-worker. You mentioned some stuff on service workers and construction workers where they're coming out with two different -- quite -- quite a bit difference in their dose reconstructions. you go a little bit further on that, please? 8 MR. RINGIN: I'm glad you bring up -- actually didn't think I'd prompted you to ask these questions, but I had -the question about extrapolating from co-workers to coworker is very, very difficult for construction workers. We've been spending years trying to come up with predictable models for exposures for workers doing similar kinds of tasks. Bob Herrick* -- who many of you know at Harvard -- has worked with us in a working group, and we spent I think two years doing -- just reviewing workers who were doing turnkey maintenance construction on cold-fired utility boilers, thinking

that that is a very similar kind of exposure. And we

went in there, he did measurements on a lot of these

so great that we couldn't draw conclusions from it.

different sites, and all we got was a variance that was

we couldn't come up with a statistically predictable model for -- for exposures. And my guess is that that's going to be somewhat similar for radiation. 4 I can also comment a little bit -- unless I'm over-extending 5 my time here -- one of the problems of the way that DOE had characterized its work sites, and I think that the beryllium example is a good one, both for Hanford and for Savannah River. The belief was that construction workers wouldn't have been exposed to beryllium because construction workers don't work with beryllium. DOE had characterized the environments where beryllium They'd done wipe samples and that kind may have been. of stuff. But after we got all these results on -- at 3 Hanford, for instance, three to four percent of the workers testing positive on the beryllium lymphocyte proliferation test, we started to go back and look at how they had been sampling and estimating environment. And in those buildings they did the wipe sampling up to eight feet on the -- on the walls. They'd not done sampling in the rafters, not above ceiling tiles, not behind wall panels or in subflooring or crawl spaces. 2 Well, that's where most construction workers do their

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work. Most construction workers don't do new
      construction. Just about all the construction work
      that's done now is either repair, maintenance,
3
      renovation, demolition or decontamination, and that's
      where they probably get most of these exposures.
      they're exposures that are very hard to predict because
      the environment isn't anticipated and the work that
      they're doing isn't anticipated. I don't know if that
      answered your question --
0 MR. ESPINOSA: Yeah, that answered my question.
1 DR. ZIEMER:
              Thank you. Other questions or comments?
                                                        Thank
      you, Knute, for your input to the Board.
3 We've come to the completion of today's agenda. We begin
      tomorrow morning again at 8:00 o'clock in terms of the
      informal time together, and then the official business
      beginning at 8:30.
7 Let me pause and see if we have any housekeeping items,
      Cori, tonight to address? No.
9 MS. HOMER: The only thing I would suggest is that if you
      have anything in the room, take it with you.
1 DR. ZIEMER: Okay. Don't leave things in this room over --
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overnight. They're -- they will become part of David

Brenner's comedy act. 2 With that, we'll recess till tomorrow morning. Thank you very much. 4 (Whereupon, an adjournment was taken to Wednesday, December 5 10, 2003 at 8:00 a.m.) DECEMBER 10, 2003

PROCEEDINGS

REGISTRATION AND WELCOME

2 DR. ZIEMER: Good morning, everyone. Again I'll call the meeting back to order, the second session of the nineteenth meeting of the Advisory Board on Radiation and Worker Health. Again I remind you if you have not registered your attendance at this meeting, please do so at the -- the books are out on the table at the entryway. Also, members of the public who wish to address the Board, please sign up in the sign-up book that also is at the entryway.

1 Again remind everyone that there are other documents and handouts on the table here to my left. Please avail

yourselves of those, as you might find it helpful. 2 Many of you are aware -- perhaps all of you are aware -that the Board has been seeking contract support for 3 assistance in auditing the dose reconstruction process and related matters. The firm that was the successful bidder was Sanford Cohen & Associates, and from that company we have this morning with us Dr. John Mauro, who's going to give us a general briefing about their company. 0 I want to point out both to those here on the -- at the podium or -- in other words, the Board members as well as those in the audience, that this presentation and discussion this morning is very general. The Board has yet to review in closed session the document which is proprietary which will be addressed this afternoon, the task orders and the independent government cost estimate for that support, so the Board will be looking at that in detail this afternoon. But basically this morning we're introducing John and the company to the Board and to the public. So John, we're pleased to have you here this morning, and if you would give us an

overview of your organization and related matters.

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1 Question?
2 MR. GRIFFON: Just a question before we start. Is -- can we
      discuss or ask John questions regarding the technical
      skill proposal or --
5 DR. ZIEMER:
              I'm going to ask Martha to address that.
      understanding now is that in fact that is still
      restricted information until the Board has discussed
      the proposal, so that that may be off-bounds.
      we'll ask Martha -- if you would, Martha -- to address
      that for us and give us a legal opinion here.
1 MS. DIMUZIO: We did speak with contracts about this, and
      basically what we have is we have a document that's
2
      before the government for its consideration, so it's
      not really a public document until a final award has
      been made. So when Dr. Mauro is giving his
      presentation, you can ask general questions about
      anything, but we cannot have any specific discussions
      about the proposal that he's submitted or the
      approaches that SC&A may be taking in -- in their
      approach to completing those tasks. That's really
      discussion for this afternoon in the closed session.
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It's a -- the document is not a public document at this

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point. It's a document before the government for
      consideration, so --
3 MR. GRIFFON:
              And SCA can't be in the closed session.
                                                         Is
      that correct?
5 MS. DIMUZIO:
               That's correct.
6 MR. GRIFFON: So we -- I mean I'm not sure how we'd
      negotiate or discuss technical scope with the
      contractor if --
9 MS. DIMUZIO: Basically what we would be developing is we
      would be developing questions that would be referred to
      SC&A for them to respond to. I mean that's how it
      would work.
3 DR. ZIEMER: And understand that this has a practical effect
      of perhaps stretching things out a little bit because -
      - but SCA then has a right, as I understand it, to have
      a certain amount of time to respond to questions that
      are raised. They are not -- we don't sit there in the
      room and ask them to respond right at that moment.
      have to develop --
0 MS. DIMUZIO: Right, within the --
1 DR. ZIEMER: -- both the technical and the cost-related
      questions and then go back to them then, is my
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understanding.
2 MS. DIMUZIO: Right, within the scope of the contract, SC&A
      has seven days to respond to those questions, so...
4 MR. ELLIOTT:
               It's also a redirection, too. If the Board
      finds in the technical proposal that the contractor is
      proposing something beyond or -- or outside of what the
      scope of the task called for, you can redirect, by
      comment. And so that -- that, you know, of itself is -
      - as well as the questions that you formulate, are --
      are confidential --
1 MS. DIMUZIO:
               Right.
               -- in nature.
2 MR. ELLIOTT:
3 MS. DIMUZIO: We can't appear to be leading -- we cannot
      appear to be leading the contractor to arrive at a
      certain point. It's more of a direct -- you know, we
      have these questions and it's -- you know, it is
      basically that negotiation back and forth, so that's
      done in closed session.
              So those are kind of the ground rules on which
9 DR. ZIEMER:
      we need to operate this morning. Let me ask, any --
      everybody on the Board understand that?
2 MR. MILLER: Excuse me, Dr. Ziemer.
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1 DR. ZIEMER: Yes, sir?
2 MR. MILLER: Could I just raise a question on this?
      there a practical way to solve this? Because the
      technical scope of what the audit is going to be is of
      significant public interest. We're not interested in
      the money side, you know, the independent government
      cost estimate part. But it seems to me if there's a
      practical way to solve this would be if the -- if NIOSH
      would make available the accepted bid proposal that was
      submitted with Sanford Cohen & Associates' original bid
      that was made to the government that was accepted, I
      presume, by the -- whatever source evaluation board you
      had, so that there's some sense about what the
3
      structure, the organization, the methods that are going
      to be used, the approaches, what the -- what the items
      are that they're even going to do. I mean to --
7 DR. ZIEMER: Well --
8 MR. MILLER: -- to -- to go into secret --
9 DR. ZIEMER:
              Let me --
0 MR. MILLER: -- and to have discussion about what those
      items are --
2 DR. ZIEMER: I understand your point, Richard --
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1 MR. MILLER: -- without any public --2 DR. ZIEMER: -- but let me -- let me point out to you that the scope of the work has been defined by the Board. That is a public document. Our determination is whether or not this company is responsive to our scope and what the costs will be to carry that work out. the scope is public. We have several tasks. been defined. We have to determine whether or not this company is in fact capable of responding to those tasks. You will hear about the organization of the company and their personnel. 2 The question on the original proposal, I don't know the legal answer to that. I -- it's a procurement issue. 3

Perhaps Martha can speak to that.

2

5 MS. DIMUZIO: We did pose this specific question to our procurement office about is there any way that the proposal as it's been submitted can be supplied to the public, and basically at this point it's through a Freedom of Information request that would be considered by the procurement office as to whether or not the document is releasable, working with Sanford Cohen to make that determination whether or not anything within

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that proposal was proprietary or so forth.
2 But I think it's important to point out that while SC&A may
      have developed an approach and everything, not that
3
      there's anything wrong with that approach, but it may
      not be the approach that the Board wants.
                                                 And so I
      think we need to, you know, have that discussion --
      that's a discussion that needs to be held in private,
      whether or not the way Sanford Cohen has determined to
      approach the work that needs to be done is the way that
      the Board intended it to be. I mean that's one of the
      issues for the session, the closed session, is to
      evaluate how their approach has --
              Is the issue of the original document, is that
3 DR. ZIEMER:
      being pursued or does that require a specific FOIA
      request?
6 MS. DIMUZIO: That would require a specific FOIA request for
      the original document, as it stands now.
              I think we were referring to the proposal that
8 DR. ZIEMER:
0 MS. DIMUZIO: Oh --
1 DR. ZIEMER: -- Cohen originally submitted, which I believe
      does contain some proprietary information.
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1 MS. DIMUZIO: As a part of the contract submittal?
2 MR. ELLIOTT: Yes.
3 DR. ZIEMER: Originally.
4 MR. ELLIOTT: I think that's what we're -- they're referring
       to.
6 MS. DIMUZIO: That would be -- yeah -- I need to go back and
       -- and speak with the procurement office, but I believe
       if Sanford Cohen is willing to release that
       information, that that information could be released,
      yes.
1 DR. ZIEMER: We will follow-up on that. You know, I don't
      want to get into that debate right now, but we -- note
       is taken of that, Richard. We'll see if it can be
       released --
5 MR. MILLER: Just to be --
6 DR. ZIEMER: -- at some point.
7 MR. MILLER: -- clear, though, that proposal that Martha was
       referring to, the original proposal, was incorporated
       by reference, I believe, in the final contract award
       that NIOSH posted.
              SANFORD COHEN & ASSOCIATES BRIEFING
2 DR. ZIEMER: Thank you. Okay. With that introduction,
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here's John.
2 DR. MAURO: Good morning. I'd like to thank you for
      inviting me.
4 DR. ZIEMER: Wait, John, you need to put it --
5 DR. MAURO: A little higher? Okay, is that a little better?
6 Again, thank you for inviting me. I'm glad to be here
      today. As was mentioned, SC&A was selected -- I guess
      it was back in October -- to provide technical support
      to the Advisory Board in fulfilling its mandate under
      the Act.
1 In November basically we were awarded what's called a task
      order proposal, which means that from time to time the
      Board would request SC&A to perform certain tasks.
3
      received our first task order request for proposal in
      November and we recently -- last week -- submitted our
      technical proposals to the Board. As they mentioned,
      they're reviewing that.
8 I'm here today primarily to introduce SC&A, who we are --
      we're a small company -- and to identi-- let you know
      our scope of work, what we've been asked to -- to do.
      And also give you the brief overview of some of the
2
      people that are on our team and their backgrounds.
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1 Okay, SC&A. We're a small company, do about $5 million a
      year in work, and we primarily -- we were incorporated
      in 1982. Sandy Cohen is a personal friend of mine.
      I've been working with him now for -- since 1986, and
      we specialize in doing dose calculations.
      nuclear engineers and health physicists, primarily.
7 We have currently 30 employees and we have 50 associates.
      We have a way of doing business whereby we have a core
8
      group of people that are more senior, like myself, and
      then we have associates that work with us who are
      specialists in a wide variety of areas that we bring in
      to work on particular problems as they arise.
      effect, we -- and this is a -- this is a very effective
3
      approach in providing technical consultant services in
      that we can bring the best people in the world to -- to
      the table to -- to answer very specific questions. And
      also it allows us not to carry a large overhead, so we
      -- what this -- it puts us in the position that we can
      bring the best people at the lowest price.
0 And we work -- we have our headquarters office in McLean,
      Virginia, but our -- a lot of our people work out of a
      virtual office. For example, I work out of my home in
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Red Bank, New Jersey.
2 Our clients currently are government clients, primarily.
      do a lot of work for the Nuclear Regulatory Commission,
      the Environmental Protection Agency, Centers for
      Disease Control, FEMA and DARPA. We -- we write
      reports. We write new reg documents. We write -- for
      EPA we write baseline risk assessments, technical
      support documents. Dose reconstructions, we've done a
      lot of work for CDC for off-site dose reconstruction.
      We've done -- so we do dose assessment, risk --
      radiological risk assessment primarily for the
      government and primarily for agencies that regulate DOE
      -- so it's, you know -- so we -- I guess one of the
      reasons for our selection is that as a corporation we
      really are not tied very closely at all to DOE. We're
      more closely tied to EPA, NRC, Defense Nuclear
      Facilities Safety Board, folks that regulate Department
      of Energy.
9 However, we do have a -- we have a laboratory, and our
      laboratory does radiological analysis of samples from
      anyone that sends us -- that wants us to do, and we are
2
      doing some work from samples that come from the State
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of New Jersey, but also from Savannah River, from -- I
      guess it's CH (Inaudible) out at Rocky Flats, so that -
      - that really is the place where we're -- we're doing
      some work for DOE through our laboratory analysis.
5 By and large, we're a consulting company to government
      agencies, and I'm proud to say now we are working for
      NIOSH, also.
8 Our organization -- we have a simple organization.
      president and owner of the company is Dr. Sanford
      Cohen. He's a Ph.D. nuclear engineer. He started the
      company in 1982, and he is a personal friend of mine.
      And he in fact will be the back-up. If for any reason
      -- I got eaten by an alligator or whatever -- Sandy is
      my back-up on this project. I'm the project manager.
5 The person -- our chief operating officer, Greg Beronja,
      he's there to make sure we make money. He -- he runs
      the operations. He has a -- he's a chemical engineer
      with an MBA.
9 But then the -- really the heart, the operation where -- the
      people in the trenches are these four boxes.
      four divisions in the company. The original company
2
      that was doing lots of work for NRC and EPA is now
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called the consulting division. I head the consulting
      division. That division, as I said earlier, consists
      of anywhere -- at any point in time, between 10 to 20
      people doing -- writing new reg documents and other
      types of paper (Inaudible) newspaper.
6 We have -- the other division is the laboratory, and they do
      radiological analysis. Their specialty is
      transuranics.
9 We have a field division, folks that go out -- and this is a
      relatively new division. They go out and perform
      measurements. If you folks are familiar with
      characterization and close-out surveys, Bill Ulicny is
      -- leads up that division and it's a -- it's just a
      start-up operation. We have a few -- a few private
      clients.
6 And finally we have what's called the quality assurance
      division, another smaller division within the company,
      and Patrick Kelly heads that up and he does audits.
      Right now he's doing some audits related to I guess
      (inaudible) whereby these waste packages are being
      produced and -- for -- for disposal and there's very
2
      formal auditing process to make sure those packages
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meet certain criteria. We drafted Patrick to help out
      a bit 'cause that's what he does, audits, and -- but --
      but all of the work that will be done in this project
      is going to be done out of my division, consulting
      division.
6 I put up the organization chart -- we'll talk about the
      individuals, and it's probably hard for you to read
      anyway. I should have made it a little larger.
      the concept of operations, the way we organize
      ourselves, is that I -- I'm the project manager and I
      will be available to this project full time, if
      necessary. I'm here at -- to serve the -- the Board.
      I have been -- I'm a key individual on the project. I
3
      cannot be replaced unless written authorized approval
      by the Board. I re-- and I report to the Board.
      Probably there's also an administrative relationship
      between the Board and NIOSH, but my understanding is
      that I will be reporting to you folks and to get all my
      direction from you folks.
0 I have a deputy project manager, Joe Fitzgerald.
      you folks may know him. He has a lot of background at
2
      the Department of Energy. He has an indep-- he has his
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own company. We brought Joe's company, Salient, in as a subcontractor to SC&A. 3 You'll see, as we go over the backgrounds of these folks here, we -- we have -- we've done a lot of off-site dose reconstruction for CDC. We've done a lot of risk assessment and dose assessment for EPA and the NRC, but we don't know, as a company, the DOE complex and the issues that you folks are dealing with. Joe brings to the table this know-how, which we consider to be very important. 1 We've broken up -- what I have -- the way I've arranged the organization is a staff that crosses -- that will be supporting me. One I call quality assurance. A lot of folks are not familiar with the concept of quality assurance versus quality control. We all come out of the nuclear industry where that's our life's blood. Ιf it is -- we have to follow very rigorous protocols to ensure quality. We have Dr. Steven* Ostrow, who is going to basically provide audits of our work to make sure that we do everything in accordance with the procedures that we have written up in our plan to you 2 folks, and he will be auditing that to make sure we do

that. 2 And to the right of me on this org. chart is our records management specialist. We'll be tal-- I'll be talking 3 about these people a little bit more when we get into each individual, but we -- our vision of the project is that it's going to be ve-- it's critical that we maintain a complete record that will be accessible to the Board. All of the information we use should be completely transparent. And I realize there's going to be a lot of hard copy and electronic versions. documents might need to be controlled, so we have a records management specialist. I'll tell you a little bit more about Kathy Behling when we get into the individuals. 5 Then we've broken up the project functionally according to the scope of work that we would be covering. We have a -- our -- a large staff to do the reviews of the individual dose reconstructions. We have a -- the center box there is to do worker and site profile reviews. And then we have a third box which is SEC petition reviews and supporting that aspect. So we've

broken up ourselves functionally into those areas.

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1 Reality is, on projects like this, it has to have a certain
      fluidity to it. All of the people that we were talking
      about are -- are available to work basically on
      whatever is needed.
5 The bottom half -- bottom half box, what we did is we went
      out and tried to find the best people we could -- this
      is where this associate relationship becomes very
      valuable -- in all of the areas that we felt were
      important to have access, ready access to powerful
      expertise to be brought in as needed to -- to get the
      job done. So we have a total of about 31 people that's
      available to the project.
3 And we -- one more point that I'd like to make is we also
      put in place a very powerful conflict of interest
      control process. We understand the importance of
      conflict of interest. And everyone that comes aboard
      and is part of our team has to go through a vetting
      process before they can be part of the team to make
      sure that we meet all conflict of interest issues.
      no one comes aboard unless you say it's okay, and we
      have a process to do that.
2 Let me just introduce -- what I've done in the org. chart is
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I've identified what I call lead individuals.
      are -- this is the heart of the operation. I'm going
      to just give you a brief biosketch on each person so
      you get to know who we are.
5 I'm the project manager. I have a Ph.D. in health physics
      from New York University Medical Center. I studied
      under Dr. Merril Eisenbud, and many of you folks may
      remember Merril. I have also been certified as a
      health physicist since -- continually since 1976, and
      my whole life has been doing dose calculations.
1 Kathleen Behling, she is our records management specialist.
       She has an associate's degree and 30 years experience
      in records management. She spent a large part of her
      career responsible for records management at GP Nuclear
      and maintaining all the occupational exposure records
      electronically and at -- in our -- at our company, and
      she's been with us now for about ten years, maybe
               She does all our records management work
      related to our dose reconstruction work for CDC.
0 As you can imagine, when you do an off-site dose
      reconstruction -- in fact I -- work we did a while back
2
      for CDC involved 65,000 boxes that we had to go through
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and -- and create electronic files, vet them out and collect electronic files, and she was responsible -together with our database management people -- to create this bibliographic database, so she's -- she's the records management specialist. 6 Hans Behling. Hans Behling has a Ph.D. in health physics and a master's in public health, 35 years experience. He spent many years at the (inaudible) after TMI to get -- straighten out the situation. He is -- as far as I'm concerned, I've been working with him now for ten years -- one of the best health physicists I've ever met, and he's a pit bull. He will dig and he will dig, and he has a great deal of experience in dose reconstruction. 5 Our company was hired by the Republic of the Marshall Islands on behalf of the claimants who were concerned that they were not getting treated right in their compensation for their claims, and we were asked to reconstruct the doses to the people of Marshall Islands from the Bravo test. The Bravo test is an infamous test that resulted in very large exposures to a large 2 number of Marshall Islanders and we were asked to come

in an independently dig through ancient records to reconstruct the doses that were experienced by the people of Marshall Islands. We came and -- our findings were very interesting. We believe the government underestimated the thyroid doses by about a factor of ten, and the whole body doses by about a factor of two, and that's very controversial -- getting a lot of heat on that, but we've got the evidence. 9 Victor Evdokimoff, Victor is a certified health physicist. His entire career was dedicated to hospital health physics. He was the -- he is recently retired as the RSO for Boston University Medical Center. And Victor is recently retired from that position and he's with SC&A now. 5 Joyce Lipsztein, Joyce is a -- has a Ph.D. in health physics. She also went through the same program that I went through at New York University Medical Center under Dr. Eisenbud, and she's I guess perhaps one of the lead-- world's leading experts on internal dosimetry. She knows IMBA like her own name. could -- she could talk the talk, so we are so pleased 2 because she also recently retired as a professor and

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has joined SC&A and she's available to us full time, so
      we have access to what I consider to be one of the
      world's experts on internal dosimetry and all of the
      software, including IMBA, that's used to reconstruct
      doses.
6 Arjun Makhijani, you folks know Arjun. Arjun is an advocate
      for worker rights and we feel that -- and -- but also
      he is a superb scientist. And so he brings to the
      table what I believe is -- how do I best say this?
      When you're in -- when you work in the nuclear
      industry, you know, sometimes your bills are being paid
      by the Nuclear Regulatory Commission or the Department
      of Energy, and what we have here is a mix of people
3
      that come from many diverse backgrounds. Arjun is a
      strong advocate for -- for worker rights. He will --
      he will lead up the SEC petition reviews, and -- and
      he's available to us -- he's one of the individuals
      that's only available to us about half-time.
      else here is available to us just about full time.
O Steve Ostrow I mentioned earlier. He's in charge of quality
      assurance. He's going to be the watchdog to make sure
      we're following our procedures and fulfilling our
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obligations under our standard operating procedures.
2 Bill Ulicny, we drafted him from our field program. He's a
      great health physicist. He's one of the younger
      members of the team and he will be a case manager.
5 Oh, one of the points I'd like to make regarding the way we
      think about this project is we are -- we believe in the
      concept of a case manager. That is, every case that we
      review, there will be a person who we -- who will be
      held accountable for making sure that case is processed
      properly. And so we have identified a number of people
      who will serve as case managers, and they have the
      freedom to draw upon all the resources of the -- of our
      project team and more, if necessary. We'll go outside
      and get whatever is necessary. So Bill will be one of
      our case managers.
6 And finally is Joe Fitzgerald. He's the president of
      Salient, a recently-formed corporation.
                                               They're a
      subcontract to us, and as you know, Joe is a -- very
      knowledgeable on the DOE complex.
0 Okay. With that, you have a pretty good idea of who SC&A is
      and the people that will be working on this project.
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now will just briefly go over the scope of work that we

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were asked to -- to write a proposal for. This came in
      back in November, and as I mentioned, on December 2nd
      we filed our proposal.
4 Task one is -- is the big task. It involves us doing
      reviews of 70 basic dose reconstructions, 70 advanced
      dose reconstructions and ten blind dose
      reconstructions. And we haven't yet received any of
      the -- I guess you would call it the administrative
      records, so how we're actually going to staff to get
      that work done will depend very much on what -- what
      the issues are and we'll staff it accordingly. And we
      will certainly keep you apprised of how we're doing
      that once -- once we get the ball rolling.
4 We've also been asked to support each of the Advisory Board
      meetings, so every one of these meetings, we'll --
      we'll be here and we will be giving briefings on what
      we found out so far.
8 We've also been asked to take a look at -- apparently there
      are procedures that have been prepared by -- by NIOSH
      to review the SEC petitions, and we've been asked to do
      a critical review of that.
2 And then of course in number six there, these are -- there's
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2

a lot of deliverables, there are lots of reports that we will be delivering. And all of this work -- this will be performed over a one-year period, once we get the green light to proceed. 5 Task two focuses in on site profiles. Basically -- and this would be -- the site profile work on this project, as I mentioned, is -- is -- would be headed up by Joe Fitzgerald, and he will be drawing upon all the resources of -- of our organization. Not only the organization chart we showed here, but whatever it takes to get the job done.

2 Basically the TORP, the task order request for proposal that we received from the Board and from NIOSH, identified a possibility that this coming year we may be asked to do a critical review of -- of 16, up to 16 site -- site profiles, and part of that work will include not only reviewing it for completeness, but also performing what we call worst-case analysis. That is, given the data, use that data to evaluate what we think the upper bound doses might have been associate with particular operations for each one of these. And that's where we bring in again our team of health physicists. So you

always think about you have the -- the radiological engineers and the health physicists looking at the site profiles, and then we have a team of specialists like Joyce who will help in evaluating what they call worstcase scenarios. A part of that work will include making visits to the sites and -- and digging and digging and digging to make sure that we turned over every stone to make sure the site profiles are as complete, that -- that -- as they can be. 0 Task three, you're probably familiar with the OCAS -- I guess it's IG-1 and IG-2. These are the procedures that are currently being used by NIOSH and their contractor to perform external dose reconstruction and internal dose reconstruction. But in addition to that, there are also -- we became aware of a large number of additional procedures that have been prepared by NIOSH contractors, and these basically are the procedures that they're following -- they're technical procedures. So this task basically involves us performing a independent technical review of those procedures. 1 We believe that the same people that are reviewing the dose 2 reconstructions should also be the people reviewing the

procedures, so they will be responsible for doing -the same people will be working on task three as are working on task one. 4 Finally our fourth and last task is called the dose 5 reconstruction review tracking. What -- what this is is that we're -- we're going to generate a great deal of information. I mean beside the records, electronic or hard copy, we receive regarding the cases or regarding the site profiles, we will be receiving -- we will be filling out -- the way we are approaching the project is a very, very formal documentation process of audits where each review follows an audit procedure, check list, sign-offs, and so we're going to be generating a lot of data and information. Our plan is to build a database management system that is compatible or integrated with your Sequel* 2000 that will allow the Board to -- to basically do sorts on -on the records, that data that's in there, that will help serve your purposes in tracking performance, doing statistical workups of the data. So -- and so -- we have Don Loomis who specializes in database management 2 system. I know -- Sequel is his -- he knows Sequel the

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way Joyce knows IMBA, so we -- we feel very comfortable
      that we can design and build for you whatever you need.
       And it's easy to do. He says don't worry about this
            The others are going to be some tough ones.
      Don't worry about this. We'll -- we'll build whatever
      you want.
7 And that -- that really concludes my overview. I'd like to
      just make one statement. I've been in -- I've been
      doing dose calculations for 30 years. This is the most
      important project I've ever been on and I'm very, very
      pleased that you selected us. Thank you.
      questions?
3 DR. ZIEMER: Thank you very much, John, for that overview of
      your company and capabilities. We'll now open the
      floor for questions from the Board. Again, I'll remind
      the Board members that you pretty much need to confine
      your questions to the material that John has presented
             That restricts your questioning right away,
      doesn't it?
0 John, you indicated that in addition to the roughly 30
      folks, you have the capability of bringing others in on
      rather short notice. Is that correct?
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1 DR. MAURO: Yeah, we've -- we've built up -- right now we have 50 active associates. We have 30 full-time employees and 50 active associates. However, we've -we develop associate -- we actually at one time had perhaps 200 associates, so we have a relationship with a -- a network that -- that -- throughout the United States of -- for example, you'll notice Art Upton is on our org. chart.

9 DR. ZIEMER: Uh-huh.

3

2

0 DR. MAURO: Well -- or Doug Boreham -- I don't know if you know Doug, he was up at (inaudible) University. specializes in biomarkers. These -- these are folks that are -- that I consider to be the best people there are out there to address particular questions. We have a list right now of hundreds of specialists in all the radiological sciences and the nuclear sciences that -that we have an ongoing relationship with. I mean all of the -- the key people -- you'll notice by the list of names here, we -- we all have 30 years experience under our belt. Collectively we have a network of relationships, of people that we can draw upon. So when special problems arise -- in fact, for example,

Roy DeHart.

before the Board -- I don't know this individual, but is a Dr. Hunt who's -- I believe Great Britain, that Joyce said listen, you -- we've got to get Dr. Hunt abo-- available because there's no one who understands film badge dosimetry and converting film badge readings to organ doses better than he does, so we brought him We envision that there are going to be aboard. problems that are going to come up that are going to be very specialized. And so what I'm saying is that what we're in a position to do is very quickly bring aboard associates within a day -- for some reason, if we have a need -- so we -- there's no boundaries. We can either do that, or we can bring aboard a subcontractor. That's more of a difficult thing to do. There are companies that have certain specialty expertise that you may want to bring in. We're very much open to doing that. But usually that takes a little longer because we have to put the contract in place, and the vetting process regarding conflict of interest becomes a little more burdensome. But -- but yes... 1 DR. ZIEMER: Okay, thank you very -- oh, here's a question.

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1 DR. DEHART: I actually was going to ask the question, but
      you answered it as you were talking through, and that
      is most of our -- essentially all of our cases that
      we're going to be reviewing have medical problems.
      They have cancer. And I noticed that there was the
      absence of any physician being listed, but when you
      mentioned Art -- you took care of that issue.
8 DR. MAURO: Okay, right. Yeah, Art and I are -- are
      friends. He -- he was part of the NYU -- New York
      University Medical Center program and over the years I
      -- we've been working with him. He's been an SC&A
      associate for many years.
3 DR. ZIEMER: Okay, thank you very much. We appreciate your
      being here today.
       REVIEW AND APPROVAL OF DRAFT MINUTES, MEETING 18
6 We now have opportunity for a working session. We have
      several items that we need to address. First of all,
      beginning with the minutes to the 18th meeting -- let
      me get my copy back from Ray here. I'd like to ask
      Board members for any additions or corrections to the
      minutes.
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2 Let me begin by indicating that in the executive summary I

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have asked Ray to insert on -- looking for a page
      number here. It's the first page of the executive
      summary. It would be the third section under OCAS
      Program Status Report, David Sundin's report where he
      announces the number of claims to date. I asked Ray to
      insert the number of claims in the summary here so that
      it is more specific in the executive summary.
8 Likewise at the top of the next page where it indicates four
      completed site profiles, to identify those four sites
      in the executive summary.
1 Now other -- other comments? And again we're asking for
      substantive ones as opposed to simply grammatical. Did
      you all check those items that are attributed to you to
      make sure -- Wanda, you have one that you wanted to
      raise, I believe.
            I have two. On page 20 where Ms. Munn begins
6 MS. MUNN:
      talking. We've just finished talking about the
      procedure for individual dose reconstruction, and it's
      not clear to me in that sentence exactly what I was
      talking about.
1 DR. ZIEMER: Let's make sure at page 20 -- page 20, I'm
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looking to make sure because I have a downloaded

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version which seems to have ended up with different
      page numbers and so on, but --
3 MS. MUNN: All right, the --
4 DR. ZIEMER: -- this is the copy that was in our packet, I
      guess.
6 MS. MUNN: Yes, in the center of the page, the motion to
      approve the procedure for processing passed
      unanimously.
9 DR. ZIEMER:
              Okay.
0 MS. MUNN: And then the next sentence, Ms. Munn indicated
      she was still concerned about the large number of site
      profiles being required.
3 DR. ZIEMER: Okay, so to -- for clarity, to add the words
      "of site profiles" --
5 MS. MUNN: "Of site profiles being required." Because our
      discussion was about how quickly these were going to be
      done.
8 DR. ZIEMER: Right. And so without objection, we'll add
      that.
0 MS. MUNN: And the other concern, page 32, second paragraph.
       Jim Neton was talking here and the first sentence of
2
      the second paragraph says "Determination of external
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doses was covered by general considerations,
      unmonitored workers." I'm not certain exactly what
      that means. I think I'd have to go back to the
      transcript to get the full sense of that, but that
      sentence appears to need some grammatical correction of
      some sort. I'm not sure exactly what.
7 DR. ZIEMER: Yes, it's -- does not appear to be clear to --
      does anyone know what -- Jim, she's attributing this to
      you.
O DR. MELIUS: No, Jim Neton.
1 DR. ZIEMER: Oh, to Jim Neton. Oh, oh, the other Jim.
      Okay. Okay, so we will ask for that sentence to be
      clarified.
4 Does that cover the ones -- okay. So we don't know how that
      will be fixed right now, but we will fix it.
6 DR. MELIUS: Just put a mumble in there. Dr. Melius
      mumbled; couldn't understand a word he was saying.
8 DR. ZIEMER: I found the Gen Roessler statement that I was
      trying to understand.
0 DR. ROESSLER: Okay.
1 DR. ZIEMER: It's on page 32. It's bullet five.
      Roessler asked what part of the total was assumed for
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the chest X-ray. What part of the total.
2 DR. ROESSLER: I think on that Jim was talking about typical
      doses to workers, and he had a chart up and he showed
      chest X-ray and I -- I kind of thought that was a maybe
      a major part of the dose and so I asked him
      specifically what -- what dose was due to the chest X-
      ray, and he didn't specifically have an answer.
      just said it would be typical for whatever was used in
      medical facilities at the time. Does that help?
O DR. ZIEMER: What part of what total? A typical total
      worker dose?
                    Is that --
2 DR. ROESSLER:
                I think that's what he was talking about, as
      best I can remember. We might have to go back to the
      slides he was using, but I seem to recall he had one
      slide where he showed a typical dose and he included a
      chest X-ray.
7 DR. ZIEMER: So it would be something like what part of the
      total in the example? Well, perhaps we can ask that --
      that that --
O DR. ROESSLER: I think we need to the record.
1 DR. ZIEMER: It's not clear to me exactly what they're
      talking about.
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1 Other -- other items that anyone wishes to call attention
            Did you have another one, Wanda, or is --
      to?
3 MS. MUNN:
           No.
4 DR. ZIEMER: Okay. No others? Let me ask for -- if the
      group is willing for us to make appropriate fixes to
      those two spots, if you then are willing to approve the
      minutes, including the executive summary, as slightly
      modified. Is there a motion to that effect?
9 MS. MUNN: So moved.
0 MR. PRESLEY: Second.
1 DR. ZIEMER: It's been moved and seconded. All in favor of
      approving the minutes, subject to relatively minor
      fixes, please say aye.
                    (Affirmative responses)
5 DR. ZIEMER: Any opposed?
                        (No responses)
7 DR. ZIEMER:
             Motion carries.
                            (Pause)
9 DR. ZIEMER:
              Larry wants to ask a question relative to
      minutes.
1 MR. ELLIOTT: These minutes are very detailed in their
      content, and I would just ask for the Board's sense on
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what you would be happy with. Is this what you're
      happy with as far as your minutes, or would you --
3 DR. ZIEMER: As far as level of detail, I think you're
      asking.
5 MR. ELLIOTT: Right. Or would you prefer perhaps to see
      something like the executive summary, six pages or
      less, and then use the transcript to rely on that, on
      what the verbatim is to what was actually discussed and
      held. How would you --
O DR. ZIEMER: Yeah, just give us some feedback on this.
      We've been trying to condense them and they're --
2 MR. ELLIOTT: Cori, am I correct that -- that this executive
      summary and then the full text of minutes that we are
      using is not -- it's something we can change. We can -
      - the Board wants to have just an executive summary
      style set of minutes, they can do that.
7 MS. HOMER: (Off microphone) We have some (Inaudible) can be
      met in a much shorter version.
9 MR. ELLIOTT: Yes.
O DR. ZIEMER:
              Jim.
1 DR. MELIUS: Yeah, I actually like this particular style, so
      I speak in favor of keeping it. I think --
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1 DR. ZIEMER: About this level of detail?
2 DR. MELIUS: About this level of detail, 'cause I think it
      is helpful to be able to not have to refer back to the
      transcripts to find what someone said at this point in
      time.
6 DR. ZIEMER: Gen Roessler?
7 DR. ROESSLER: I think we owe this amount of detail to the
      people who are looking at our minutes, and I know there
      are people -- particularly in health physics -- who are
      reviewing what we're doing, and I think we need at
      least this much detail.
2 DR. ZIEMER: Okay. Mark, did you also have a comment?
3 MR. GRIFFON: Just the -- the same comment as Jim, that not
      -- not too many people are going to turn to the
      transcripts, so I think this level of detail is good.
6 DR. ZIEMER: Okay. Other -- any others want to weigh in one
      way or the other? Do you want them shorter, longer,
      this feel --
9 MR. PRESLEY: I agree.
O DR. ZIEMER: Seems to be a sort of a general consensus that
      maybe we're at about the right level. Okay.
                                                    Thank
2
      you, that's very helpful.
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1 Okay, Ray, I think -2 THE COURT REPORTER: Who made the second to approve them? I
3 didn't hear -4 MS. MUNN: Presley.
5 DR. ZIEMER: I'm sorry?
6 THE COURT REPORTER: Who made the second to approve these
7 minutes? I didn't see it.
8 DR. ZIEMER: Who made the second to approve the minutes?
9 Okay. Robert, thank you.

BOARD DISCUSSION/WORKING SESSION

(Pause)

meeting -- and I'll simply remind the Board of it and
then we can handle it as you see fit -- was whether or
not we should have a -- a subcom-- put in place a
subcommittee, as opposed to a working group, a
chartered subcommittee to handle the ongoing issues
relating to dose reconstruction and our interactions
with the contractor. It may be that the Board will
wish to delay that decision on establishing a working - or a subcommittee until after we have a chance to
review the contractor's proposal later today. But

nonetheless, let me ask if the Board does at this time wish to move forward on that issue or -- in the absence of that, we remain operating as a committee of the whole on the issue of how we direct and work with our contractor. I'll open that for any comments or specific recommendations. Begin with Dr. Melius. I agree, it's difficult to talk about this with 7 DR. MELIUS: specificity now until we've gone into closed session. 8 However, I think -- I am a little uncomfortable about us making major changes in Board procedure or sort of the issue of the public's access to what we're -- to our activities and so forth -- to do that in closed session. So I think it would be worthwhile having some discussion of the concept and -- of subcommittee and at least some sense of where we --6 DR. ZIEMER: Yeah. 7 DR. MELIUS: -- where we should go with it and what the sub-- a subcommittee or working group might -- might do or not, but recognizing that we may have to sort of develop the specific charge for the subcommittee in closed session, given the situation, that all we can do 2 right now is --

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1 DR. ZIEMER: No, actually this could not be part of our
      closed session today.
3 DR. MELIUS: Okay, that was actually --
4 DR. ZIEMER: I'm sorry if I suggested that. I suggested
      that the decision on doing that may need to wait till
      the results of the closed session are known to us.
      we -- we would not -- we cannot carry out other
      business in the closed session other than reviewing and
      addressing the cost proposal that is before us.
      of necessity, must be an open session item. So -- and
      all I'm saying is that unless we choose to do something
      this morning on that, we would defer and would continue
      to act as a committee of the whole until our next
      meeting, at which point we could decide to establish a
      subcommittee on an ongoing basis. There's not a
      necessity that we have a subcommittee at the moment.
7 DR. MELIUS: Yeah, can I just say two things to address
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that? And some of this is going to be a question for

Larry. One is that I think first of all we ought to

look and -- and see what -- what we might lose or gain,

timing of task orders and so forth if we -- I think our

depending on what happens this afternoon, in terms of

next meeting's what, about two months away -- until then 'cause -- I mean I think if -- to the extent we could facilitate this moving forward through a subcommittee, I think it would be -- be good. It may be that we would then -- so that's one question. The second question is I think it would be worthwhile having a larger discussion of what a subcommittee might do on an ongoing basis. We may have to defer the decision on that to the -- to the next meeting. But I wouldn't see -- there certainly might be some value to having a subcommittee that would last until the next meeting and would have a very specific charge to it in terms of what it -- it might do, though I'm not sure what that charge would be until I understand the process, and even then I'm not sure we can do anything in this session on a contingent basis --7 DR. ZIEMER: Yeah, a subcommittee that lasts until our next meeting looks much more like a workgroup as it's ad hoc and very specific. But unless -- right now we would be operating in the absence of having a precise knowledge

2 DR. MELIUS: Well, I think we could -- well, I'm not sure

of what the charge would be to such a workgroup.

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'cause I'm not sure what the procedure is. I would
      think that, though -- that a -- as I understand it, a
      subcommittee can take actions on behalf of the Board
      between meetings; a workgroup cannot.
5 DR. ZIEMER:
              If the Board so authorizes.
6 DR. MELIUS: Correct. Correct. And so I guess my question
      is, number one, in the short term -- and this question
      I think is to Larry -- given the process, given what
      might occur this afternoon -- you know, what are the --
      what are the possibilities this afternoon, and then do
      any of those possibilities...
2 MR. ELLIOTT: I think --
3 DR. MELIUS: -- would be assisted by having action by the
      next meeting.
5 MR. ELLIOTT: I think it would be beneficial for the Board
      to hear again the process from -- from this point
      forward, and that'll give you a better sense of, you
      know, what kind of a delay might occur and how you
      might react to -- to that. So I'd ask Martha if she
      would again cover the ground of how -- how this thing
      is going to -- going to work.
2 MS. DIMUZIO: Well, basically, you know, we'll go into
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closed session this afternoon to review the proposals and -- and the cost estimates that have been provided by Sanford Cohen. The Board at that point can either determine to accept the proposals as -- as -- as they've been submitted, and then we would move forward with award. If there are questions related to the -the approach or level of effort that may -- the contractor may be proposing and things like that and there are specific questions, we would generate those questions, those questions would be forwarded to our procurement office, who would then provide them to Sanford Cohen for response. At that point Sanford Cohen has seven days to respond to those questions and potentially re-propose against those four tasks, and that point in time we would have those proposals to forward out back to the Board for them to approve. 7 MR. ELLIOTT: And you can put forth an extension of time if -- if appropriate and necessary and justified. 9 MS. DIMUZIO: Right, yes, if Sanford Cohen needed additional time to prepare their responses or whatever, then yes, I mean we can give them additional time of seven days to respond, yes.

1 DR. ZIEMER: Okay, let's go ahead and get some other comments here and then we'll proceed. Let's see, Henry and then Wanda.

4 DR. ANDERSON: Yeah, my -- I guess my thoughts are if the intent is to have a subgroup so that we can move expeditiously, I think we could also continue with a --I mean for a subgroup you still have to -- or subcommittee, you still have to post it in the Federal Re-- or you know, you've got to have all the advance notice and all that kind of thing. It would seem to me we could simply do that as a committee of the whole, recognize that the only thing we have to have is a quorum and, you know, if we have to have multiple calls, you know, getting a subcommittee together can probably be more problematic than just getting a quorum from the -- from the Board to address whatever needs to be talked about, but as far as the logistics of doing the announcement, it's not much different, so that might be the way -- and then as activities go, we could see whether there's some more routine activities that a subcommittee would be more advantageous, but I -- I just think we're early on enough that everybody's going

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to want to see it and probably be involved in the
      decision process. And we just have to recognize on
      short notice some people won't make it and we'll just
3
      have to be sure that, whatever the call is, we've got
      the quorum.
6 DR. ZIEMER: Wanda.
7 DR. ANDERSON: You can't do it by --
8 MR. ELLIOTT:
               By call.
9 DR. ANDERSON: You can't do it by call?
O MR. ELLIOTT:
               No. You cannot hold a closed session --
1 DR. ANDERSON: No, no --
2 MR. ELLIOTT: -- by phone call.
3 DR. ANDERSON: Okay, this would have to be a closed session.
4 MR. ELLIOTT:
               Yes.
5 DR. ANDERSON: Oh, okay, I was thinking --
6 MR. ELLIOTT: This is a negotiation --
7 DR. ANDERSON: -- a subcommittee action would be --
8 MR. ELLIOTT:
               This is a negotiation between --
9 DR. ANDERSON: Oh, this is for --
0 MR. ELLIOTT: -- you as the government and your contractor.
1 DR. ANDERSON: Yeah, okay, I see. I thought there would be
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other issues to look at. Sorry, never mind.

2 MS. DIMUZIO: That's correct.

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1 MR. ELLIOTT: Sorry.
2 DR. ZIEMER: It would be equivalent to doing what we're
      doing this afternoon with a -- some sort of perhaps a -
      - assuming we needed revisions.
5 DR. ANDERSON: Yeah.
6 DR. ZIEMER: Maybe we don't.
7 DR. ANDERSON:
               Yeah.
8 DR. ZIEMER: But if we did... Wanda.
9 MS. MUNN: Although it appears cumbersome to act as a
      committee of the whole, in the absence of a triggering
      event or substances that would -- circumstances that
      would clearly require the more concentrated efforts of
      a subcommittee, I see no reason for us to further
      discuss establishing one at this time.
5 DR. ZIEMER: Jim, did you have an additional comment?
6 DR. MELIUS: I have an additional comment. If I followed
      you, Martha, correctly -- and I'm guessing at times and
      given holiday seasons, but my sense is that the
      earliest go back and forth with Sanford Cohen &
      Associates would -- end with us getting a new proposal
      around the first of the year.
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1 DR. MELIUS: Yeah, so it would be about right. Our
      meeting's the first week in February, so we would be
      losing a month of work in terms of -- of tasks, should
      they have to be revised, et cetera.
5 I guess another question is -- and I don't know that the
      answer is -- can the process go on more than once?
      we end up going back and forth with them...
8 MS. DIMUZIO: Yes, technically you could. I mean if -- if
      level of effort was still not correct, if, you know,
      they still didn't fully understand the -- you know, how
      we wanted them to revise if necessary, then yes, you --
      you could be going back with -- with a follow-up, yes.
3 MR. ELLIOTT: Martha, can there be back-and-forth in written
      form between the contractor and the full Board?
5 MS. DIMUZIO:
               I believe so. I'd want to --
               So for example, on a specific --
6 MR. ELLIOTT:
7 MS. DIMUZIO:
               -- verify that with --
               -- task, if -- if there were --
8 MR. ELLIOTT:
9 MS. DIMUZIO:
               -- Larry, but --
0 MR. ELLIOTT:
               -- limited questions or issues regarding a
      proposal on a given task, and the Board puts that
      together and goes back to the contractor and then they
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get a revised proposal and it looks okay, the Board
      could take action on that and make an award by letter
      to procurement.
4 MS. DIMUZIO: Yes, you could do that. You would just have
      to have some mechanism for all of the Board to approve
      the questions --
7 MR. ELLIOTT: Approve and agree.
8 MS. DIMUZIO: -- and review the questions and okay the
      questions in some type of session that would not be
      open to the public. But yeah, I mean -- and I don't
      know how you would --
2 MR. ELLIOTT: Cori, is there a way to do that by -- by mail?
3 MS. HOMER: (Off microphone) I'm sorry, I was -- the music
      is a little loud back here.
5 MR. ELLIOTT: Okay. I'm trying to get at can the Board
      conduct some of this business of awarding a task
      without having a face-to-face meeting and going back
      and forth between them and their contractor by -- by
      mail, perhaps, and get a sense that -- that the full
      Board is in agreement, has a -- has a -- no?
1 MS. HOMER: I don't believe so. I -- I could check into
      that, but the -- any action taken by the Board is
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- considered a meeting, period -- has to be announced.
- If it's closed, it has to be announced as closed and
- approval has to be gained for that.
- 4 DR. ZIEMER: Yeah. Actually that sounds true. I mean it
- sounds -- it sounds like if you tried to do things by
- letter, you're circumventing the intent of the -- the
- 7 process so that --
- 8 MS. HOMER: (Off microphone) Yeah, the same -- the same
- 9 (Inaudible) for a subcommittee, as well.
- O DR. ZIEMER: I might add -- and let me ask this -- or
- perhaps make it a comment and ask a question. If --
- we're presuming that there might be changes, but maybe
- there won't. But let's assume that out of today's
- 4 session the Board raises some comments and asks for
- feedback.
- 6 MS. HOMER: Uh-huh.
- 7 DR. ZIEMER: And the result is some sort of a revised
- 8 proposal around the first of the year, as was
- 9 suggested. If at that point the Board felt that there
- was some urgency in acting on that new proposal, would
- it not be possible to announce in the Federal Register
- that in two weeks or something we're going to meet --

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1 MS. HOMER: Uh-huh.
2 DR. ZIEMER: -- I don't know, Cincinnati or somewhere for
      the express purpose, in closed session, of making the
      final -- taking the final action on that?
5 MS. HOMER: That's possible.
6 DR. ZIEMER: I know it's not optimal.
7 MS. HOMER: No, it's not. We have to have seven days.
8 DR. ZIEMER: But if there's some reason to shorten --
      otherwise we have the month, quote, loss, but if the
      Board felt that we can't afford to sit here for a month
      with nothing happening --
2 MS. HOMER: Uh-huh.
3 DR. ZIEMER: -- we need to go forward, we could meet.
4 MS. HOMER: It would be tough.
5 DR. ZIEMER: I mean whether it was the Board or the
      subcommittee, you have to do the same thing.
7 MS. HOMER: Yeah, I mean --
8 DR. ZIEMER: You have to make the announcement --
9 MS. HOMER: -- given the appropriate resources, we could --
     we could pull that off. I mean the -- the --
1 DR. ZIEMER: And could we not --
2 MS. HOMER: -- Federal Register notice --
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1 DR. ZIEMER: -- even reserve some time in advance for that
     possibility?
3 MS. HOMER: I would suggest we do that, yes.
4 DR. ZIEMER: And then if we didn't need to use -- use the --
      it could -- is it easier to cancel an --
6 MS. HOMER: It's very difficult to cancel --
7 DR. ZIEMER: -- announced meeting than it is to add one?
8 MS. HOMER: -- hotel arrangements.
9 MR. ELLIOTT: I think --
0 MS. HOMER: If I have a contract with a hotel, it's very
      difficult to can-- I'm sorry.
2 MR. ELLIOTT: I think for the benefit of this discussion,
      though, we -- you could have your closed session and --
      in the offices --
5 MS. HOMER: Yes, we could.
6 MR. ELLIOTT: -- at NIOSH in Cincinnati. And let's be
      minimal in what the expectations are to set up that
      meeting.
9 MS. HOMER: Yes.
0 MR. ELLIOTT: That would be a seven-day advance notice for
      the Federal Register that we'd have to put in.
2 MS. HOMER: We have to have seven days to get it published.
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It can be published on an emergency basis.
      Determination to close can probably be rushed. As long
      as we are not dealing with a contract with a hotel, I
3
      don't foresee a problem with that. I mean rooms are --
      I'm sure we can find lodging for you at a hotel which -
      - which would not --
7 DR. ZIEMER:
             Larry volunteered --
8 MS. HOMER:
              -- mean a contract.
             -- four bedrooms.
9 DR. ZIEMER:
O DR. MELIUS:
              I can't wait.
1 DR. ZIEMER:
              I don't want to be in --
2 MR. ELLIOTT: I have a tent, too.
3 DR. ZIEMER:
              Jim.
4 DR. MELIUS:
             To me, rather than doing something contingent
      on a full Board, I think let's explore the subcommittee
              We've got some time this morning and seems to
      issue.
      me that if a subcommittee could be charged -- well, I
      guess this is the question again. Can the subcommittee
      be charged with, you know, reviewing a response from
      the contractor should be -- should it be necessary,
0
      based on our meeting this afternoon, and then approving
2
      it? And -- and the instructions for that approval, the
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circum-- you know, whatever you want to call --
2 MR. ELLIOTT: With the bounds on it.
3 DR. MELIUS: -- with the bounds on it could be, to some
      extent, set this afternoon by -- by what our
      instructions are back to the -- I mean I think in
      essence we end up doing that when we -- the questions
      and -- should we send those back to the contractor. I
      mean to me it would be a lot more --
9 DR. ZIEMER: What you're proposing if you're bounding it,
      for example, in terms of dollar values, I don't think
      we can do that in -- at this point.
2 MR. ELLIOTT: You couldn't do that in this meeting.
3 DR. ZIEMER: In this meeting.
4 MR. ELLIOTT: You couldn't do that in this public meeting.
      You could -- you could do that in -- if you set up a
      subcommittee or if you set up, as part of your
      discussion in the closed session this afternoon, what
      your expectations are and the next meeting is either a
      subcommittee or a quorum of this body, that's your
      guidance.
1 MS. HOMER: I would like to --
2 DR. MELIUS: And that's going to --
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1 MS. HOMER: I'm sorry, Dr. Melius, but I would like to point
      out that you can make the decision to establish a
      subcommittee, but administratively it still has to be
      established prior to any meeting taking place.
      Whatever --
6 DR. ZIEMER: Has to go through the --
7 MS. HOMER: -- decision that you make --
8 DR. ZIEMER: -- CDC process.
9 MS. HOMER: Correct, uh-huh.
0 MR. ELLIOTT: It has to have a charter.
1 MS. HOMER:
             Yes.
2 MR. ELLIOTT:
               The charter has to be signed off on.
3 MS. HOMER: Well, it's an establishment memo that will
      provide membership, it will provide the function, it
      will provide frequency of meetings --
6 MR. ELLIOTT: Delegation of authority from the Board.
7 MS. HOMER: Well, and that's something -- yeah, we would
      probably have to discuss that.
9 DR. MELIUS: How long?
0 MS. HOMER: I suspect it could take two weeks. With the
      holidays, maybe a little longer.
2 DR. MELIUS: Christmas Eve we'll...
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1 DR. ZIEMER: Comment over here.
              If I understand, we would need only a quorum of
2 DR. DEHART:
      this committee, which is 50 percent plus one, and the
      odds of being able to find that number to attend a
      meeting is probably greater than having a subcommittee
      with limited numbers to be able to meet, so I see no
      advantage at all at this point to try to create and
      generate a subcommittee and just have the -- just have
      us as a whole try to address the issue.
O DR. ZIEMER: Okay. Other comments?
1 DR. MELIUS: Yeah, I guess I was just thinking the opposite,
      that -- that we could establish a subcommittee of say
      four people and those four people could pick a date a
      lot easier than whatever the quorum is -- what, six or
      seven -- I don't even remember what a quorum is, seven
             One more than half.
7 DR. ZIEMER:
              -- seven, so seven -- four people are easier to
8 DR. MELIUS:
                        The question is, can we come -- I
      meet than seven.
      think the real question would be more do we want to
      spend the time and can we come to agreement at this
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meeting on a subcommittee charter, or is that going to

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be something that's going to take us more than one
      meeting to work out.
3 DR. ZIEMER: Well, it's not obvious that four is easier than
      seven because it's a specific four versus any seven out
      of 12, so I'm not sure -- it's not obvious to me that -
      - if it's a specific four, it may be actually harder to
      find a date, but that's what you're suggesting.
      who knows, it depends on who the four are and who --
      what the dates are, so who knows.
0 Other comments? Henry.
1 DR. ANDERSON:
               I guess the other thing to talk about, what
      would be the activities going forward of such a -- I
      think it's advantageous to have a relatively small
3
      group which would basically act as our project officer
      for the Board, as a collective group dealing, so kind
      of moving forward it would seem to me if there were
      questions that had to be dealt with --
             Well, the original idea on this was more of a
8 DR. ZIEMER:
      management type --
0 DR. ANDERSON: Yeah.
1 DR. ZIEMER: -- of group that would --
2 DR. ANDERSON: Yeah.
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1 DR. ZIEMER: -- work with the contractor closely, help
      decide who --
3 DR. ANDERSON: Yeah.
4 DR. ZIEMER: -- which of the Board members would participate
      in different cases --
6 DR. ANDERSON: Yeah.
7 DR. ZIEMER: -- and so on, as opposed to a specific decision
      such as this one --
9 DR. ANDERSON: Right.
0 DR. ZIEMER: -- which is on the contract itself. Okay.
      Other -- a comment, Cori?
2 MS. HOMER: Yes, also something to consider would be that
      even if a subcommittee has been formed and the
      establishment has taken place, the Board still has to
      meet to determine what authority they're going to give
      to the subcommittee. That's still going to take a full
      meeting of the Board, at least one.
8 DR. MELIUS: Again, just a question. Wouldn't you do that -
      - couldn't -- if we did that today, doesn't that
      establish the charter and the -- the --
1 MS. HOMER: It does establish the charter, but it does not
      establish the authority that the Board is going to give
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the subcommittee. Because the way that it works,
      without any authority, the subcommittee cannot take
      action without approval of the full Board.
4 DR. MELIUS:
             But --
5 MS. HOMER: If the Board decides to give the subcommittee
      authority to act on their behalf, that authority has to
      be developed and approved before it can -- you know,
      before the subcommittee can take any action.
9 DR. ZIEMER: Can that authority be given prior to the
      approval of a charter, which is what --
1 MS. HOMER: I would suggest not.
2 DR. ZIEMER: -- which is the situation that we would have
      today.
4 MS. HOMER: You could develop them at the same time.
5 DR. ZIEMER: But until the charter was approved, the
      authority could not --
7 MS. HOMER: The authority has no -- no -- you have no
      authority.
9 DR. ZIEMER: This is a little knotty, but I think in the
      interest of moving forward -- and we can return to this
      -- I think I'm simply going to rule that we will
      continue to operate as a committee of the whole for now
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and -- particularly on this issue, and if necessary,
      try to establish some kind of a -- what I might call an
      emergency meeting of the Board if we need to do
      something before our next meeting. In the meantime, a
      -- more details on the charter can be developed.
      actually there is some work that's been going on.
                                                        Mark
      and I have worked together on some draft things for a
      possible charter and -- and may be that we'll have a
      chance to present that a little later even today, Mark,
      for this subcommittee.
1 We do need to take a break and then return, so let's take a
      break till 10:15.
3 (Whereupon, a recess was taken.)
      ADMINISTRATIVE HOUSEKEEPING AND BOARD WORK SCHEDULE
5 DR. ZIEMER:
              This is the time on our schedule that we take
      care of some administrative/housekeeping issues.
                                                        We'll
      first turn the mike over to Cori to see if she has any
      particular items that she needs to bring to us.
9 MS. HOMER: (Off microphone) Just a few things. Am I on?
0 Okay. A couple of things just real quickly. Please don't
      forget to e-mail Larry your work time for the Board,
      prep time, any working groups that you -- that you were
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on or worked for, and your Board time separately. Go
      ahead and send that to Larry as soon as you can, and cc
      me.
4 I wanted to mention, I have some trouble getting all of the
      e-mails quickly. That delays payment for everybody, so
      please respond as quickly as you can with your time to
      Larry so that I can get you guys paid quickly.
8 The closed session will be held in the Mesquite I Room, just
      past the front desk, down that long hallway.
      first room on your left-hand side. It's taking place
      as scheduled on the agenda.
2 DR. ZIEMER: At 2:00 o'clock at Mesquite --
3 MS. HOMER: At 2:00 o'clock at Mesquite I. Now for those
      that are going on the tour, we have a very, very full
      agenda for that day. Please dress casually.
      will be no cameras, phones, blackberries, palm pilots,
      no forms of communication allowed, period.
8 MR. PRESLEY:
               Picture I.D.
9 MS. HOMER: Uh-huh. Bring water, because there will be ice
      chests, if you'd care to. And we'll be departing
      around 6:15, 6:30. I have an agenda and I will make
2
      you copies, but I haven't had the opportunity to do
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that just yet.
2 DR. ZIEMER: So you want people in the lobby at 6:15. Is
     that what you're saying or --
4 MS. HOMER: Yes. My apologies, but that's the agenda. They
      arrive at about 6:15 with the bus.
6 DR. ZIEMER: Yeah. And it's a drive out to the test site,
     an hour and a half, roughly.
8 MS. HOMER: Well, they're saying about an hour, hour and 15
      minutes.
0 DR. ZIEMER: Okay.
1 MS. HOMER: Please be ready to pay --
2 DR. ZIEMER: So what -- what's the time to meet in the
      lobby, very speci--
4 MS. HOMER: Around 6:15, 6:30 --
5 DR. ZIEMER: Around? Exactly --
6 MR. ELLIOTT: Give it --
7 MS. HOMER: 6:20.
8 DR. ZIEMER:
             6:20.
9 MS. HOMER: How's that? That'll give the bus five minutes.
0 MR. ELLIOTT: How about if you say this. The bus leaves at
     6:30.
2 MS. HOMER: There you go, the bus leaves at 6:30.
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1 MR. ELLIOTT: Be there or --
2 MS. HOMER: Or miss the bus.
3 MR. ELLIOTT: -- miss the bus.
4 MS. HOMER:
            Since we've ordered lunches, for those that have
      ordered lunches, please be prepared to pay -- I believe
      it's $6.95 -- to their guide. I'm not going to be
      collecting any cash, so --
8 MR. PRESLEY: And they do take -- that's all they do take is
      cash.
0 MS. HOMER: Is cash, so -- any questions?
1 DR. ZIEMER: It can be in quarters, if necessary.
2 MS. HOMER: Yeah.
3 DR. ZIEMER: Nickels, for the big spenders.
4 MS. HOMER: Period, they will confiscate them.
5 MR. ELLIOTT: Not on the bus.
6 DR. ZIEMER:
              What --
7 MS. HOMER:
             That tell -- he was -- David was asking if we
      could even bring anything on the bus, and we cannot.
      Leave everything in your room.
0 DR. ZIEMER: And by everything, you're talking about --
1 MS. HOMER: I'm talking about the electronics.
2 DR. ZIEMER: -- any cameras, any --
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1 MS. HOMER: Yes.
2 DR. ZIEMER: -- electronic things.
3 MS. HOMER: Uh-huh.
4 DR. ZIEMER: Don't even bring them there.
5 MS. HOMER:
             That's right. They'll be confiscated.
6 DR. ZIEMER: Permanently.
7 MS. HOMER: No, they'll be given them back at the end of the
      tour when we arrive back at the hotel. That's --
      that's what I've been instructed.
0 DR. ZIEMER: Thank you.
1 MS. HOMER:
             Will that be it?
2 DR. ZIEMER: Any questions for Cori?
3 MR. ELLIOTT: When you turn your time in to me, keep in mind
      that the working group on the Board audit or the --
      Mark Griffon's working group has now completed its
      charge and as of yesterday I believe the working group
      on -- Dr. Melius's working group on evaluating the
      interview process had completed its charge, so working
      groups have a finite life and they have -- once they
      meet their charge, then we can't bill time against
0
      them.
2 DR. ZIEMER: But the time that they spent up till now --
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1 MR. ELLIOTT: Yes, yes.
2 DR. ZIEMER: -- but don't come in two months or three months
      from now --
4 MR. ELLIOTT: Well, the point here is the working group on -
      - Mark Griffon's working group on DR evaluation
      essentially completed its charge last meeting, so
      between last meeting and this meeting there shouldn't
      have been any -- any effort for that.
9 MS. HOMER: Okay. Now did you want to discuss dates for a
      potential --
1 DR. ZIEMER: Yes, that's in fact the next item.
2 MS. HOMER:
             Okay.
3 DR. ZIEMER: Let me remind you that we have -- we are slated
      for the 5th and 6th of February, the meeting site being
      Augusta. And we may want to identify a contingency
      site in early to mid-January. This is -- we do require
      I think the presence of the Federal official, so we
      need as a front end -- I know that Larry's schedule I
      understand is pretty busy in January and so Larry, are
      there -- I think we start with you and I guess we also
      are required to have a Chair.
2 MR. ELLIOTT: I think so.
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1 DR. ZIEMER: But my guess is that your schedule in January
      is probably tighter than mine. I have decided not to
      attend the Citrus Bowl game on January 1st where Purdue
      will play Georgia, and my only conflict right now is
      the 30th of January, so I'm okay. Where are you,
      Larry, on --
7 MR. ELLIOTT: January 12th, 13th and 14th I plan to be in
      Richland with the site profile team disseminating that
      bit of information out there in -- for the Hanford
      folks, so I'll be on travel then.
1 DR. ANDERSON: (Off microphone) Are you going to be there
      for the health effects subcommittee meeting, too?
3 MR. ELLIOTT: I hadn't planned on that, don't know anything
      about that, so...
5 DR. ANDERSON: 'Cause that's the 22nd and 23rd, I think.
      No, no, it's the 15th and 16th.
7 MR. ELLIOTT: No, I had -- I had not planned to participate
      in the health effects subcommittee.
9 Then on the -- January 2nd of course wold be not a good day.
0 MS. HOMER:
1 MR. ELLIOTT: That's coming back from the holiday. And the
      19th would not be a good day, either. That's a --
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- that's a holiday I'd kind of like to take this time.
- 2 And then the -- looks like the 26th and 27th would not
- 3 be good days.
- 4 DR. ZIEMER: Let's start with the week of the 5th then --
- 5 MS. HOMER: Dr. Ziemer, I have leave that week.
- 6 DR. ZIEMER: Okay, that takes care of that week. Let's look
- at the week of the 12th. So you're out the 12th, 13th,
- 8 14th --
- 9 MR. ELLIOTT: And 14th.
- O DR. ZIEMER: -- and is the 15th a travel day also, then?
- 1 MR. ELLIOTT: No, I'll be coming back on the 14th.
- 2 DR. ZIEMER: We actually -- if we need a, quote, emergency
- meeting, we're talking about a one-day maximum, I
- 4 think, so let -- let me ask about the 15th and 16th,
- 5 are either of those days bad for anyone?
- 6 DR. MELIUS: 16th is a -- may be problematic for me. The
- 7 15th works, though.
- 8 MR. GRIFFON: 15th is okay.
- 9 DR. ZIEMER: Okay, 15th is a possibility? Let's look at the
- o next week, 19th?
- 1 MS. HOMER: No, somebody couldn't.
- 2 DR. ZIEMER: 19th is a holiday.

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1 DR. MELIUS: Yeah, the week of the 19th I'm tied up all
      week.
3 DR. ZIEMER: Tied up all week. Others? And remember, we
      actually only need a quorum, but if possible, we'd like
      to get everybody there -- 20th through the 23rd are
      days then when we -- is there any -- are there more
      than one person not available on those days, and just
      make a note --
9 MR. PRESLEY: I can't be there the 22nd or the 23rd, but I
      can be there the 20th and the 21st.
1 MR. ESPINOSA:
               I'm out the 23rd.
2 DR. ZIEMER: Once we're beyond that, we're almost up to our
      other meeting, so there's no point in going further.
4 DR. ANDERSON: After the 19th, we might as well put it off.
5 DR. ZIEMER: Right. As -- there are possibilities, if
      necessary, 20 and 21, but it looks like perhaps the
      15th might be the day then. Should we go ahead and
      block that out? Is that too soon after you get back?
9 MR. ELLIOTT: Huh-uh, that's --
O DR. ZIEMER:
              You're all right? Is that agreeable?
      kind of half-way between --
2 MS. HOMER: The 15th is fine for me.
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1 DR. ZIEMER: It's fine for counsel or -- okay.
2 MS. MUNN: It's fine for me. I can fly back with Larry on
      the 14th.
4 DR. ZIEMER: Okay, then let's set aside Thursday the 15th as
      a special meeting, if needed.
6 MR. PRESLEY: Cincinnati?
7 DR. MELIUS: Are we talking Cincinnati?
8 MS. HOMER: I think that would be best.
9 DR. ZIEMER: Cincinnati's all right?
0 DR. MELIUS: Can we do it like 11:00 to 2:00 or something so
      people can fly in --
2 DR. ZIEMER: Fly in and fly out? Sure.
3 DR. MELIUS: -- day trip, I think it would also allow I
      think the --
5 MS. HOMER: Will that be enough time?
6 DR. MELIUS: -- west coast people to get back out.
7 DR. ZIEMER: 11:00 to 2:00 or 11:00 to 3:00?
8 MR. ELLIOTT: Yeah, we can do that.
9 MS. HOMER:
             Okay.
0 DR. MELIUS: And that would avoid the hotel issue, to some
      extent.
2 MS. HOMER: Yes.
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1 MS. MUNN: Only to some extent.
2 DR. MELIUS: Where are we going to put --
3 DR. ZIEMER: Or it would minimize the --
4 MS. HOMER: The Westin.
5 DR. ZIEMER: -- overnights, right. And that would be at
      NIOSH.
7 Okay, is that agreeable? Any objections?
                        (No responses)
9 DR. ZIEMER: Okay. So pencil that in. Does the Board wish
      at this time to also look ahead into the March/April
      time frame and set aside some dates?
2 MS. HOMER: We already have, Dr. Ziemer, for April.
3 MR. PRESLEY: I was going to say, we set aside --
4 DR. ZIEMER: That's right --
5 MS. HOMER: April in Richland.
6 DR. ZIEMER: Yes, we --
7 MS. MUNN: We have 20, 21, 22.
8 DR. ZIEMER: I see it, it's here. Actually we set aside 19
      through 23, did we -- did we finalize the dates?
0 MS. HOMER: No, I haven't.
1 DR. ZIEMER: Okay. We sort of --
2 MS. HOMER: Yeah, just keep that week open and --
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1 DR. ZIEMER: So that -- well, I think -- settling on it was
      going to be dependent on what you could find --
3 MS. HOMER: What was available, yes. Wanda and I are
      working on that right now.
5 DR. ZIEMER: Okay. So that basically takes us up to May.
      That covers the next six months then.
7 Question arose as to the possibility of a Savannah River
      Site tour. Bob, can you help us -- whether -- how that
      would be done? Is that something you can help with?
0 MR. PRESLEY: Yes. Yeah, the guy that used to be the head
      of Savannah River is now at Oak Ridge heading up DOE so
      if y'all want to go to Savannah River, I can call him
      when I get back and see about a tour, if that's -- if
      that's what everybody wants, that's up to y'all.
5 DR. ZIEMER: Are you okay on that, Ray? I wasn't sure it
      was -- okay.
7 Let me -- let's get a straw vote here. Board members, how
      many of you would like to tour the Savannah River Site,
      show of hands quickly.
0 MR. PRESLEY: Now it would have to be on Wednesday the 4th.
       The tour would have to be before the meeting because
2
      we're going to meet the 5th and 6th, so it would have
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to be on a week day, the 4th.
2 DR. ZIEMER: Show of hands, Savannah River Site tour?
                    (Affirmative responses)
4 DR. ZIEMER: One, two, three, four, I would go if you had it
      -- I've been there a few times, but you can never see
      it all, actually.
7 MR. PRESLEY: How many?
8 DR. ZIEMER: Looks like five.
9 MR. PRESLEY: Six, sev-- how many people on the staff?
O DR. ZIEMER: Looks like another five or so.
1 MR. PRESLEY: So we're talking about 10 to 12 people.
2 DR. ZIEMER: Again, this would be restricted. Right?
      Spouses --
4 MR. PRESLEY: Yes, sir.
5 DR. ZIEMER: -- could not attend?
6 MR. PRESLEY: Don't know about spouses down there yet.
7 MS. MUNN: Nice boat trip.
8 DR. ZIEMER:
             You can check on that, perhaps.
9 MR. PRESLEY:
              Yes.
0 DR. ZIEMER: Okay. Okay, Cori?
1 MS. HOMER: Absolutely.
2 DR. MELIUS: Can I bring up one other issue regarding the
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meetings? I think the suggestion was made yesterday in
      public comment period that we have availability
      sessions or, Larry, you want to call them a public
      comment session in the evening to accommodate people's
      work schedule and so forth, can that be arranged for I
      guess the next two meetings that are on the schedule?
7 MS. MUNN:
            That's rough. That's rough, Jim.
              I would think it's sort of the Board's call if
8 DR. ZIEMER:
      you want to meet in the evening, is it not? I mean --
      now we couldn't have done it here 'cause Dave Brenner
      probably wouldn't give up the auditorium. But if we
      can find a spot...
3 MS. MUNN: Well --
4 DR. ZIEMER: Do the Board members object? And we could
      adjust the meetings so that we didn't go all day and
      all evening, if you wanted to do that.
7 MR. ELLIOTT: We may be -- I'd like to hear the sense of the
      Board, but we may -- as you think about this, we may be
      limited in our ability for space. Once we -- we have
      to contract this with --
1 DR. ZIEMER: With the hotel.
2 MR. ELLIOTT: -- with the hotel, and given their
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availability...

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2 DR. ZIEMER: We could look into that. Also during the public comment period today, if any of those who 3 comment from the public might provide the Board their views on whether you think this would be valuable for members of the public to be able to attend in an evening session as opposed to during the day, why we'd be glad to solicit that input, too. We heard from a speaker yesterday that suggested that that might be a useful thing.

1 There was one other thing suggested yesterday that had to do with making known -- outside of the perhaps the official routes of publishing in the Federal Register -- to local people the presence of this Board. example, it appears that there's not a crowd of Nevada people here at this meeting. And one would say well, what was the advantage then of coming to Nevada if in fact no one from this area attends? And one of the issues is how well do the -- particularly the potential claimants and workers in the area know that we really are here, so we need to be thinking perhaps about how -- are there some other channels to develop that

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information besides the official channels that are
      being used in the Federal Register, and I know there's
      a big e-mail list that -- and so on.
4 Now obviously we can't drag people in off -- well, we could
      drag people in off the street, but -- but at least make
      sure that certain people know. I mean I -- they may
      not come anyway, but -- can we think about how that
      might -- I don't know if you've had a chance since
      yesterday and maybe the staff --
0 MS. HOMER: Dr. Ziemer --
1 DR. ZIEMER:
              -- can think about ways that might be utilized
3 MS. HOMER: -- if I may... previous experience with other
      committees, and there have been other methods used to
      announce meetings. For example, a meeting announcement
      might be prepared and distributed to -- to various news
      agencies or newspapers, TV stations, things of that
               I'm not sure exactly how we might go about
      doing that, but it's certainly been done in the past.
      I would have to also check into what our options might
      be in other areas, what -- what methods are being
2
      pursued by other committees right now.
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1 DR. ZIEMER: We had a nice turnout in St. Louis, but I think
      you'd have to say that that's largely due to the work -
      - the effort of the local person. Denise made special
      efforts to get people out. Maybe there are folks at
      other sites that might be key contact persons that
      might be helpful. I don't know if others have some
      ideas that might help here that --
8 DR. MELIUS: Can I just comment -- point out that -- not
      necessarily through any fault of NIOSH staff, but this
      hotel's had an active picket line up for the last six
      months by the building trades and therefore I don't
      think anybody from the building trades, which are most
      of the people we're talking about Nevada Test Site
      would come into this hotel to appear, and that's a
      major -- a major issue.
6 DR. ZIEMER: Right. So is there some mechanism for
      identifying those kind of issues in advance?
                                                    I'm not -
9 MS. HOMER: Well, when I contact the hotels to make
      arrangements in the future, I can always ask them if
      they are a unionized hotel or if there are any other
      union issues that I need to be aware of. But I have to
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rely on the information that they provide me with. 2 DR. ZIEMER: Okay, thank you. 3 DR. MELIUS: I've had discussion with Larry. I think we can provide some additional information. You just happened to pick the one non-union hotel left in Las Vegas, at least near the strip, so -- probably why it was available. 8 MS. HOMER: Well, I wasn't aware. Okay. Now any other general comments, 9 DR. ZIEMER: housekeeping nature, any other items we need to address? Wanda? Thank you. 2 MS. MUNN: I guess one thing I would request in our efforts to better inform the public of our meetings, it would 3 be most helpful I think if in our meeting announcements we mention what we do so that people do not have the mistaken notion that we are an adjudicating body or that we hear individual claims, because I think it's misleading for people to think they may have an opportunity to be speaking to people who will have a

bearing on how their claim is viewed, when our

responsibility is one of process, not of individual

claims. It would be very nice to have more public

attention to what we are doing, but at the same time I don't think it's fair if we don't make it very clear to them what it is we are doing.

A good point. On the other hand, we need to 4 DR. ZIEMER: recognize that the process is within the framework of the individual claims, and sometimes a knowledge of what's happened in individual claims cases helps us understand where the process may or may not be working.

This is true, no question. 9 MS. MUNN:

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O DR. ZIEMER: I think we all recognize that when we hear people relating particular stories, that we are not in a position to act on that particular case, but we may in the process learn something either about the site, about how claims are being handled and processed. in that sense, I wouldn't want to discourage people who wish to come to the Board with -- and I think most of those who would come to talk to us, with some exceptions, general public individuals, truly in the general public, are people who have claims or are either directly or on behalf of a relative, and who may have concerns that they think that might help us in the process. So in that sense, we don't want to discourage

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that.
2 MS. MUNN: No, that wasn't my intent. No. No.
3 DR. ZIEMER: Right. Yes, Henry.
                I mean one -- one thing that might help is if
4 DR. ANDERSON:
      we were to put out and say we're very interested in
      hearing about people's experiences with the process and
      their -- you know, the interviews and -- and paperwork.
       I mean we heard about paperwork, but that might be the
      kind of thing to -- 'cause I would assume the people
      who are interested in coming are those that have filed
      claims and they want to indicate when they submitted it
      and how it's been processed. And I think feedback to
      us as to how --
4 DR. ZIEMER: Yeah.
5 DR. ANDERSON: -- how --
6 DR. ZIEMER: And that's the --
7 DR. ANDERSON: -- even though it's going to be a biased
      sample, I think it is helpful to get a sense of what
      their perception is and things like that.
O DR. ZIEMER: Yes. Other comments or input on that or other
      related... And I think, Board members, if any of you
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      have particular ideas on -- or can identify individuals
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or groups that might be useful to contact at a given
      location, I'm sure the staff would welcome that.
      -- sometimes it's just a matter -- we can provide the
      information, but who do we send it to?
                                               So if we can
      help identify those, that would be useful.
                                                  I assume
      that's the case, Larry?
7 MR. ELLIOTT: Yes, absolutely.
8 DR. MELIUS: And if it would -- helpful, at least for me,
      Cori, if you would let us know as soon as possible
      about the availability of a room for the evening,
      'cause that makes some difference in terms of how you
      outreach to people.
3 MS. HOMER: Well, normally the room is available on 24-hour
      hold by contract. That's how I always set it up.
                                                         The
      -- having an evening session will also have to be
      announced in the Federal Register.
7 DR. MELIUS: Right.
8 MS. HOMER:
             So that will have to be identified ahead of
      time, as well. I don't foresee that there'll be any
      problem having an evening session, as long as we know
      about it in advance and can announce it.
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2 DR. ZIEMER: And in fact an evening session could be simply

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a time devoted for public comment, if necessary.
      don't have to conduct other business necessarily.
3 Okay, are there any other items dealing with the Board work
      schedule or administrative that need to be addressed at
      this time?
6 DR. MELIUS: I have one follow-up to yesterday 'cause I'm
      not sure what the -- what the plan is. There was the
      Congressional letter from Quinn, Slaughter and Reynolds
      to the Advisory Board and --
0 DR. ZIEMER: Right, and that letter was addressed to me.
                                                           As
      I indicated, I did study it on the plane coming out
             I want to discuss with the Department and
      perhaps with legal counsel -- some of the things
      suggested in the letter appear to me to be well
      outside the charter of this committee, but I need to
      identify that and I need to prepare a response to -- to
      those Senators.
8 DR. MELIUS: Actually Congressmen, but --
9 DR. ZIEMER:
              Yeah, they were -- three Congressmen, I'm
      sorry. That's quite correct.
1 DR. MELIUS: Can that be shared with the committee then so
      we --
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1 DR. ZIEMER: I'd be glad to do --
2 DR. MELIUS: -- can understand what's going --
3 DR. ZIEMER: I'd be glad to do that, and I think the
      committee has received copies of that, so you can --
      you can reflect back to me if you have particular
      comments to me on that. But I will prepare a letter
      and that will be made available.
               BOARD DISCUSSION/WORKING SESSION
9 The next item on our agenda is further time for discussion
      and working session. I think we completed all the
      items before us. Is there any other item that needs
      discussion at this time? 'Cause if there is not, I'm
      going to suggest that we go ahead with public comment,
      but --
5 DR. MELIUS: Could we talk a little bit about the agenda for
      the next meeting then?
7 DR. ZIEMER: We could certainly talk about agenda.
      would be quite in order.
9 DR. MELIUS:
              And I --
O DR. ZIEMER: In fact -- yeah, let's kick that off.
      another workgroup that you're involved with, Jim, and
2
      maybe you could suggest --
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with --

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1 DR. MELIUS: I'd say the research group should have a report
      at the next meeting. I think Henry and I will figure
      out our schedules finally and maybe on Christmas Day
      we'll both be home in the office and do that, so
      research group's --
6 MR. ELLIOTT: (Inaudible) Russ?
7 DR. MELIUS: Russ, yeah.
8 DR. ZIEMER: So the research subcommittee -- certainly be on
      that agenda.
O DR. MELIUS:
              I had suggested yesterday that -- requested
      that Jim Neton or someone give us a presentation on how
      the site profiles are being used in the individual dose
      reconstructions, just to walk us through some of that.
       I think that -- that would --
5 DR. ZIEMER: Sounds good.
6 DR. MELIUS: At least for me that would be helpful. I don't
      know if others...
8 MS. MUNN:
           Yes, I agree.
9 MR. ELLIOTT:
              I'm assuming there you would like to see
      examples of dose reconstruction conducted under the
      site-wide document, under site-specific type documents
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1 DR. ZIEMER: Maybe both.
2 MR. ELLIOTT: I'm sure both --
3 DR. MELIUS: Both, both, yeah, yeah.
4 MR. ELLIOTT: -- and see kind of a sampling of those.
5 DR. MELIUS: A sampling of those, but I think -- I'm going
      to leave it up to your discretion, but it would be I
      think helpful to look at -- particularly on a -- what I
      call a complex site, like Savannah River, how that's
      being used, and then some of the others, with then
      examples, you know, appropriately masked and so forth
      from, you know, individuals and what's being done.
2 MR. ELLIOTT:
               Okay.
3 DR. ZIEMER: Okay, good. Other agenda items that anyone
      wishes to identify at this time for that meeting?
5 DR. MELIUS: Can we work on -- between now and -- some sort
      of way of getting moving forward on this possible
      subcommittee issue, how to --
              The answer is -- the answer is yes, and I think
8 DR. ZIEMER:
      I'll -- I'll commit -- and Mark and I have done some --
0 DR. MELIUS: Okay, yeah --
1 DR. ZIEMER: -- some work on that and -- and we'll prepare a
      -- I think a straw man document for us. I've also
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asked Cori to provide me with the details on exactly
      what it takes to set up a subcommittee in terms of the
      structure and the ground rules.
4 DR. MELIUS: Yeah.
5 DR. ZIEMER: So we'll provide all of that.
6 DR. MELIUS: Good.
7 DR. ZIEMER: So --
8 MR. ELLIOTT: That could be held under a general agenda item
      of -- like we have Board discussion and working
      session, so I won't -- I'll just leave it at that.
1 DR. ZIEMER: But make sure -- well, we need to make sure
      that that's earmarked --
3 MR. ELLIOTT: You want a specific agenda item earmarked for
      that?
5 DR. ZIEMER: Let's earmark it so it doesn't fall between the
      cracks.
7 MR. ELLIOTT: So that's --
8 DR. ZIEMER: That will remind us --
9 MR. ELLIOTT: -- a discussion on subcommittee...
              This'll be a -- for lack of an exact title
O DR. ZIEMER:
      right now -- subcommittee on dose reconstruction.
      Actually it's broader than that, but --
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1 DR. MELIUS:
              Yeah.
2 MR. ELLIOTT: Next meeting --
3 DR. ZIEMER: Subcommittee on dose reconstruction reviews.
4 MR. ELLIOTT: At some point in time in the near future,
      probably your next meeting, you're going to have to
      identify, from the pool of completed cases, those that
      meet your -- your sample --
8 DR. ZIEMER:
              Right.
9 MR. ELLIOTT: -- for assignment.
O DR. ZIEMER:
             Right.
1 MR. GRIFFON:
               I'm not sure I...
2 DR. ZIEMER:
              Question?
               I mean -- yeah, yeah, we -- there's -- there's
3 MR. GRIFFON:
      quite a few things we probably need to do or the
      subcommittee needs to do. I don't know that the
      individual reviews, if there's a large enough pool to
      sample a lot from right now, but -- but there's also
      site profile things and -- and other things we can get
      rolling on.
O DR. MELIUS: My recollection is that Pete Turcic is supposed
      to talk next time and -- about the outreach that
2
      they're doing?
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1 MR. ELLIOTT: That is correct, he's made that commitment.
              Okay. Would it -- would a presentation from
2 DR. MELIUS:
      NIOSH on your outreach sort of complement that and --
      is that -- does that make -- make sense as a --
5 DR. ZIEMER: Put outreach on the budget (sic). Pete will be
      there representing Labor, and if there's some
      complementary things that NIOSH could say -- okay,
      here's what we're doing that complements what Labor is
      doing, it would be useful.
0 Any other items?
                        (No responses)
2 If in the interim something jumps into your mind that you
      think we ought to place on the agenda, you can let
      Larry know or you can let me know, because we'll
      develop the agenda jointly and -- and then you'll have
      another opportunity to see the draft agenda.
                                                    It can
      always be modified.
8 DR. MELIUS: Maybe Ted Katz can present the final SEC regs
      to -- next time.
O DR. ZIEMER: Am I not correct that the -- well, I won't ask
      the question.
2 DR. ANDERSON: That was rhetorical.
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1 DR. ZIEMER:
              Yeah.
2 MR. GRIFFON: Paul, is there a possibility of -- since by
      next meeting we're -- all Board members are going to
      have a copy of the IMBA software, could we have a
      training session at night? This doesn't have to be on
      Board time or whatever, but is it possible to arrange -
8 MR. ELLIOTT: Like we did for NOCTAS?
9 MR. GRIFFON: Yeah, yeah, like a training session.
O DR. ZIEMER: Well, one of the questions will be can you have
      a training session at the location of that meeting, and
      that -- it might be that the training session would
      more easily be done at the special meeting in
      Cincinnati.
5 MR. GRIFFON: Yeah, that's true.
              So we might -- we might, if it's possible at
6 DR. ZIEMER:
      that time, have some of the Board members -- 'cause we
      can't -- we're not going to do it all at once as a full
      Board, but training sessions for individuals who might
      be available prior or after that meeting, as a
      possibility. They can -- they can look into it and let
2
      us know.
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1 MR. ELLIOTT: I would offer this. At any point in time a
      Board member's traveling through Cincinnati and they
      want to stop by, they want to have an afternoon
      available to them, we'll give you some training.
      Right, David?
6 MR. SUNDIN: I think I've been volunteered.
7 DR. ZIEMER: Provided somebody's there to train us. Thank
      you.
                     PUBLIC COMMENT PERIOD
0 We're a little early for the public comment period, but if
      the commenters are here -- and I think they are -- and
      are willing to proceed, first Denise Brock is here from
      the St. Louis area. And Denise, welcome and please
      bring us your comments.
5 MS. BROCK: Hi, I'm Denise Brock, for the record. I have
      several questions today, as usual, and some comments
      I'd like to raise with the Board.
8 First of all, I would again like to say thank you for coming
      to St. Louis. We really appreciated that.
0 We've had a chance to look at the TBD for the Destrehan
      Street site, it's been a little bit of time since
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you've been there, and I would also like to know if

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there has been any further discussion with the Board
      about coming back to St. Louis to address that with the
      claimants. Perhaps we could get a bigger meeting place
      and maybe more of a crowd this time and have time for
      Q&A?
6 MR. ELLIOTT: The Board has not had that discussion, but at
      NIOSH there has been a discussion and a -- a plan of --
      being developed to go around and, as we've done at
      Savannah River Site last month, as we're getting ready
      to do at Hanford next -- next month. I can't say today
      where we're at with regard to the schedule of -- of
      those site visits, but yes, we do intend to come back
      to St. Louis and -- and talk about Mallinckrodt.
4 MS. BROCK: Wonderful. If -- if there would perhaps be any
      scheduling conflicts in the near future, if you would
      consider coming to St. Louis, I would be more than
      happy to -- to do whatever I could to help you draw in
      a crowd. I don't have a problem doing that, usually.
9 MR. ELLIOTT: We will certainly be contacting you, Denise.
0 MS. BROCK: Thank you, Larry. I also understand that there
      have been some more Mallinckrodt Chemical worker
      records that have surfaced in Georgia. Are you aware
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of that or is that something -- I understood that it
      was just something that happened rather recently and
      that DOE managed to get ahold of those. Are you aware
      of that?
5 DR. ZIEMER: We'll ask Richard Toohey if he can respond to
      that question.
7 DR. TOOHEY: I'm not aware of DOE coming up with anything,
      but on one of our data capture trips to the Atlanta
8
      National Archives or Federal Record Center, whatever it
      is, as we routinely go searching through boxes looking
      for such, we did find some more files on Mallinckrodt.
       But I honestly don't know what they contained, but
      they -- they are certainly being reviewed, analyzed and
3
      would be incorporated into the Technical Basis Document
      if there was anything in there that we didn't already
      know.
7 MR. ELLIOTT: We're not aware of anything that DOE has
      provided on Mallinckrodt at this point, so --
9 MS. BROCK: And perhaps --
0 MR. ELLIOTT: -- the discovery's been at -- at the benefit
      of our labor and ORAU's labor.
2 DR. ZIEMER: Give credit where credit is due here.
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1 MS. BROCK: And perhaps that's right, maybe that was a
      misunderstanding on my part. If I understand
      correctly, could that possibly be then internal -- or
      actual individual data on people or -- or site
      information or both, or you just --
6 DR. TOOHEY: I don't know.
7 MS. BROCK: You don't know, that's quite all right.
8 DR. ZIEMER: I'm sure as that becomes available, it will be
      incorporated --
0 MS. BROCK: It will be added to the TBD then as we -- we go
      forward.
                Thank you.
2 And in reference to the Weldon Spring and Hematite
      facilities, claimants have noticed and commented on the
3
      fact that in NIOSH or ORAU correspondence that neither
      are listed as up and coming site profiles. Is there an
      expected time line on either one of those? And if so,
      could you give me an idea of that that is? And we're
      also somewhat curious, because they are all
      Mallinckrodt facilities -- I understand that obviously
      they were different materials that they were working
      with, but I'm a bit perplexed as to why they don't all
2
      get grouped into one, why the TBDs for that one
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facility are not all done at once. 2 DR. TOOHEY: Dick Toohey, ORAU, for the record. Okay, good questions. The -- we did, as a result of the St. Louis meeting, move Weldon Springs up on the list for site profile production, and we expect to be starting on that shortly after the first of the year. 7 The other one, the Hematite facility, I -- I don't think that's on the drawing board. Unfor -- I don't have the list with me, unfortunately, but as always, we try to be responsive to the Board or -- or the public's interest and we can certainly move that up, and especially since, as you mentioned, sites did much the same thing, it should not be too hard to do. 4 DR. ZIEMER: Thank you. 5 MS. BROCK: And that brings me to my next question or comment, and I think Dr. Toohey -- yeah, just stay; I'm sorry -- Dr. Toohey and I had spoke about this earlier. I've had several situations in where claimants are coming to me with a problem. A lot of times these workers worked not just at one facility, but perhaps two or even sometimes three. And the problem arises --2 first of all, when perhaps there's a miscommunication

somehow or the records are missing, just -- perhaps as Mrs. Ehlmann spoke yesterday, her husband had worked at two facilities and for some reason, even within the 3 same Department, she's getting different stories. that causes a situation where perhaps if a worker was at the downtown St. Louis site and they are being dose reconstructed as we speak, perhaps that person could be dose reconstructed and compensable before having to wait for further TBDs to even be completed and so I was hoping that perhaps ORAU could take a look at any of the workers or claimants that have worked at more than one facility. And if in fact they were at the downtown site, could they be dose reconstructed just to see if they meet that compensability? 5 DR. ZIEMER: Dr. Toohey? Simple answer: 6 DR. TOOHEY: Yes. 7 DR. ZIEMER: Thank you. 8 MS. BROCK: And for the record, I wanted to know if all of

the phone interviews in the dose reconstruction for the Destrehan employees -- and that is just for the employees that worked at Destrehan or the St. Louis site -- are those -- are those all completed?

1 DR. TOOHEY: I wouldn't say they've all been completed, but as far as I know, all the ones have been completed where the -- we feel the files are ready to move into dose reconstruction. Some of them where people had worked both downtown and also Weldon Springs we kind of put on hold because we didn't have the Weldon Springs site profile done. Others, as we reviewed the files, we find some inconsistency or other problem, say the ICD-9 code doesn't match the cancer diagnosis or the -the employment dates seem to be inconsistent or something like that, and we look at those and try to get that corrected before we actually queue them up for interview and dose reconstruction. 4 DR. ZIEMER: Thank you. Further questions or comments, Denise? 6 MS. BROCK: Sorry, just a few more. And actually I don't --I don't think any more for Dr. Toohey. One thing that I noticed, and maybe I didn't correctly understand what I read, but I noticed that on the Technical Basis Document or actually on a dose reconstruction, I guess -- it's what it was, it was a dose reconstruction, the

Dupree-Ellis -- Elizabeth Dupree-Ellis was cited, and

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that just alarms me for the simple reason -- I guess
      this is a comment -- that she completely excluded
      internal dose, and I felt like she grossly
      underestimated things. I just wanted to make that
      comment.
6 And I don't know if I can do this. I wanted to ask a
      question of Dr. John -- is it Moreau?
8 UNIDENTIFIED: Mauro.
             Mauro? And I don't know -- can I ask it?
9 MS. BROCK:
      guess you won't necessarily be able to --
1 DR. ZIEMER: Depends on the question.
2 MS. BROCK: Okay, can I say it to the Board, say yea or nay,
      either one?
4 In reference to off-site exposure, Hematite, Weldon Spring
      and St. Louis I understand had a lot of residual
      radioactivity or ground water problems, air problems,
      things such as that. And I was curious if a member of
      the public could request Sanford Cohen & Associates to
      look into that? Like there's a situation in Hematite
      where there's residual radiation. The Department of
      Energy doesn't want the responsibility to clean this up
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      because they're saying there was a problem with nuclear
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subs. Everybody passes the buck and so nobody wants to
      clean up this mess. And I'm wondering how that can be
      addressed or how would I go about that and if that's
      something that they're able to handle.
5 DR. ZIEMER: It seems to -- it seems to me that in terms of
      their role with the Board that that could certainly be
      inappropriate, but that's something the legal folks
      would have to address.
9 MR. ELLIOTT: As a member of the public, you're -- the
      contract with Sanford Cohen & Associates is with the
      Board as a government entity, and they're given a
      specific charge, a specific scope of work that they're
      proposing against. And what you're asking for is not
      included in that scope.
5 MS. BROCK: So find other health physicists. Right?
      Basically.
7 MR. ELLIOTT: And you'd have to have the money to support
      that -- that effort.
9 MS. BROCK: Right, or maybe attorneys would, I'm assuming,
      because I think that's what a situation was in another
      area, too.
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2 Let's see. And another comment was, in reference to dose

reconstruction, in or with the absence of datas (sic), I understand that there is to be the surrogate coworker data and the use of site profiles are extrapolation. My concern is, again, how do you know that these datas are -- you're using are not -- are even accurate? There is a distinct probability -- at least in the case of Mallinckrodt -- that there was altering or coverup of datas or numbers, if you will. And again, back to those badges, I -- I understand that somebody was referencing to badges; I think that was Dr. John -- I'm sorry, Morau, Moreau --

2 UNIDENTIFIED: (Inaudible)

2

3 MS. BROCK: -- yeah, okay, Mauro, and I -- I've heard repeatedly from workers that these badges were useless, and it just is a grave concern of mine that when somebody's looking at a badge and you have two workers working side by side and one comes -- comes up red hot and the other comes up with a big fat nothing, and when you go in there to try to dose reconstruct these people and that badge is coming up with no reading, that is a big concern to me and I'm really afraid that when you use those sort of readings, I don't know how you can be

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so sure that the datas are accurate, and I guess that's
      my comment. Thank you.
              Thank you, Denise. Let me ask if any Board
3 DR. ZIEMER:
      members have questions. Yes, Jim?
5 DR. MELIUS: I have a follow-up question -- I think it's for
      Dick -- but based on what Denise asked, though. But is
      the -- for the site profiles, is there a schedule of
      those site profiles, a sort of a listing --
9 DR. TOOHEY:
              Yes.
0 DR. MELIUS: -- on the web site with an estimated --
1 DR. TOOHEY: It's not on the web site, but we supplied it to
      NIOSH, so I --
3 DR. MELIUS: Would it be possible to put on --
4 DR. TOOHEY: -- could get it --
5 DR. MELIUS: -- my -- I guess my question is would it be
      possible to put that on the web site so that people
      would, you know, have a way of knowing sort of what the
      schedule is and -- and you know, possi-- it doesn't --
      doesn't have to be, you know, exact, but estimated
      spring of whatever, something like that or --
1 MR. ELLIOTT: We'll --
2 DR. MELIUS: -- (Inaudible) subparts are.
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1 MR. ELLIOTT: We'll consider that. I can't give a guarantee today, but we'll consider that. This is a -- this is a plan that's being reviewed and evaluated right now for its feasibility and its realisticness -- if that's a word -- realism, I guess -- can we achieve it, so we're working toward that.

7 DR. MELIUS: No, and I think if it -- once it's -- the plan is finalized, that's the point I think it might be 8 helpful for the public to be able to see, you know, what you're -- you're planning on doing before it gets all...

2 DR. TOOHEY: Can I make another comment, possibly partial answer to Denise's last one on accuracy of coworker data or badge readings or anything? I would just like to mention -- don't forget, every dose reconstruction we do contains an estimate of the uncertainty in that value. And many times that uncertainty can be very large. And then that gets run through the IREP program in the uncertainty on the probability of causation. And since we're at the 99 percent confidence interval for the decision criteria, a lot of the errors or inaccuracy in the point estimates that we get in the

workers are accounted for by including the uncertainty in those values.

And many people don't realize that in most 3 DR. ZIEMER: cases, larger uncertainties help the claimant because it spreads that distribution out more. It's one of the exercises I have my students do. They -- I give them some hypothetical problems to solve with IREP and to look at the effect of uncertainty on the award or on the probability of causation. Generally it tends to favor the claimant, the more uncertain that information is.

2 Our next speaker from the public is Richard Miller.

Richard, welcome. Richard's with the Government Accountability Project.

5 MR. MILLER: Richard Miller, for the record. Good morning. I guess with respect to the schedule and agenda, I'd like to offer a plea for you to enjoy the lake effect off of Buffalo at some point. I think that there's enough interest in what's going on in western New York that I know there's probably going to be some other efforts to communicate with people up there. But there seems to be an awful lot of interest in the work of

NIOSH and the Advisory Board, and I think -notwithstanding the delightful climate there, I think that might be worth looking at as a potential future location. There's certainly plenty of people I think would be happy to cooperate with the Board and NIOSH and in having either evening sessions or outreach activity to ensure a full and robust participation, given that it has one of the largest concentrations of facilities in the country. 0 Second suggestion is -- is -- well, let me just -- as a footnote to that, I think it would be helpful to have some discussion about whether and if the Board or what policy the Board will address if people want to get a site profile reviewed. You -- Dr. Melius raised one letter that just came in from three members of Congress, and Dr. Ziemer said something to the effect of well, these are legal issues and I'll respond. I -- I think there's a broader question here, which is you've got -- as you develop these site profiles and in those sites where there's some, you know, concentrated community interest -- certainly like St. Louis -- it 2 wouldn't surprise me if you'll see more than one of

these come forward because the audit process is in fact the only check and balance on this program in -- in -within the program's boundaries itself. I mean there's other checks and balances in government, but this is the -- the design of the program. And so, you know, I was sort of looking at -- at John Mauro's presentation and there was -- I saw there was like ten to 12, you know, DOE sites and four -- two to four AWEs, I'm thinking two to four AWEs, hmmm, well, what happens if you get six letters from six Congressional areas, you know, or districts or facilities wanting you to do site profile reviews? What do you say to them? Too busy, come back next year? Legally we can't take your request? I mean I don't know what the answer's going to be, but it seems to me there's -- there's probably an opportunity here for some -- for some policy development about what the Board takes in in terms of these kinds of inputs and then how do they get resolved and addressed -- or prioritized, for that matter? mean you could spend all your time looking at uranium facilities and -- and miss Hanford and Savannah River in your site profile reviews, and so part of it may be

an allocation of resource issues. But I think that -but on the other hand, I think there's this intense public interest and obviously Congress is an important stakeholder in all of this -- this program, as well. And I just -- I just think that given, you know, this -- this -- the program is now ripening to the point where the audit function's going to start to take on, you know, flesh and bones and a real activity, that it would be very helpful -- 'cause I don't think this is something that ought to be dressed simply as a legal question. I think it's a policy question at large, so that's just my suggestion to the Board. You all think about how you want to address that. I have my own thoughts on that, but I don't know that this is the time to do it. 6 The third question is -- has to do with Blockson Chemical. This is now the third time I've raised Blockson Chemical and I know that NIOSH and the Secretary of Health and Human Services haven't asked for the Board's opinion on Blockson Chemical, but I'm going to see if I can't spur you all to kind of stick your nose into this

a little bit, 'cause it's a really interesting what I

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think is policy question. Again, I don't think it's a
      legal question alone. And that has to do with whether
      or not you include, and how much of this chain of
      production you include, of radon exposure at these
      uranium phosphate facilities that were used for -- I
      mean -- I mean rock phosphate facilities that were used
      for uranium extraction.
8 And the -- the -- oh, dear, I apologize. And I guess the
      thing that -- the thing that is interesting is --
      excuse me, I thought I turned that off -- that's
      somebody else's. I don't know.
2 That -- the Blockson Chemical issue -- I don't know, has
      anybody had a chance or -- just to the extent you have
3
      had a chance to look at the Blockson Chemical report,
      you'll see in there there's a section on radon which
      says reserved, so -- and -- and claimants have been in
      touch with me and I've had the pleasure of chatting
      with a number of people in the Joliet, Illinois area
      who have received their dose reconstruction reports.
      They have not been sent to DOL for adjudication.
      They're being held in abeyance at this point. And the
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      question is, where along the food chain of this -- of
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the intake of the rock phosphate through to the production of phosphoric acid and through the various precipitating processes and oxidation processes and through to the final uranium process and -- and so forth, do you or do you not include radon exposure? And I am aware that there are a variety and a diverse set of views because just like Gaul, this program is divided into three Federal agencies and -- and here I think each of the agencies may even have their own views on this subject. But it is as much a policy call about where you draw the boundaries around what constitutes attributable radiation exposure for purposes of this program as it is whatever some lawyers decide to concoct. And the reason I say that is, having had a chance to review the contract between the Atomic Energy Commission and Blockson Chemical, they purchased it by the pound. And so they wound up with a purchase of all of the inputs from the raw phos-- rock phosphate coming in from Florida all the way through until the uranium was extracted. And so the economic transaction would argue for a broad encompassing approach to including the radon exposure. And there

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are other ways of slicing this would say but wait a
      minute, that rock phosphate also came in and was used
      to make trisodium phosphate, Tide detergent, and should
      the program be compensating people for making Tide.
      Right? I mean you could -- you could make that
      argument.
7 And on the other perspective, one could say well, maybe
      there's a middle ground to be carved out here and --
      and we can find some sort of discrete spot to carve it
      out because this was a multi-purpose facility. You
      could also argue that this facility's economic life was
      substantially extended and bolstered because they ma--
      they did this uranium extraction, and it was a very
      lucrative contract for Blockson Chemical. And -- and
      but for this contract existed, that facility existed in
      -- in economically turbulent times when the fertilizer
      industry was definitely on the bottom of the commodity
      cycle. So there's a lot of ways to debate this
                 I'm not here to present every single one of
      those choice points.
1 I guess all I'm floating is that I think this is an item
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      which is also ripe for Board consideration. Now that
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doesn't mean that NIOSH is inviting your comment or even interested in your comment. But at the end of the day, I don't see how the Board can not look at this kind of question, because at the end of the day, you're going to be auditing it and these are generic issues to numerous phosphate to uranium processes. This isn't the only plant, as many of you know. So I just thought I would add that to your list of items where a deliberative process -- you could review the engineering reports. NIOSH I think has done an elegant job of laying out in the site profile the production process, so you could at least see it, maybe see the source documents, the contracts, and begin to grapple with this to figure out where do you draw the line. Otherwise, I think that -- that this is going to get decided behind closed doors. You're going to be left auditing something where it's going to be preordained whether or not you get to even examine that dose because the agencies have prejudged it. And I think what -- there's an ongoing deliberative process now. don't think -- I don't know, Larry, maybe you can --I'm prepared to stand corrected here, but I don't

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believe that issue is fully closed, at least as of last
      week, so I just thought I'd lay that out as a -- a
      topic where I think your expertise in combination here
      would be really valuable.
5 Am I to prepare to stand corrected, Larry?
6 MR. ELLIOTT: I have no comment.
7 MR. MILLER: Well, that gives you a flavor, unfortunately.
8 Next item I'd like to put on the plate for the Board, I
      guess -- again, and -- and for NIOSH, it's just a
      suggestion that a policy be developed with respect to
      professional standards -- conflict of interest I quess
      is sometimes too narrow a term, but it -- we've all
      used it in shorthand here to cover the broad
      professional standards of -- of -- of -- that are
      expected here, that go well beyond financial conflict.
       And -- and -- and I just can't help but observe that
      it seems like almost every meeting we have kind of a
      dandelion coming up in the lawn of another conflict of
      interest sprouting. And it seems like it's largely
      attributable and -- and frankly, fortunately, that
      NIOSH was willing to provide some transparency on who's
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      working on these teams, but people seem to be
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identifying these things for you, and it seems to me this ought to be a Federal agency function up front. They ought to be combing through this issue up front and the public serves as a check and balance, rather than the public serves as the only policeman for these professional standards questions. And so I would just urge you all to -- to think very hard about whether you want to revisit this. As Jim Melius noted at the -yesterday's meeting, in a way, a lot of us sort of thought the site profiles would accumulate the value of the knowledge of each individual dose reconstruction, and so the site profile and the conflict of interest issue really didn't ripen until -- as this program ripened. And I think it's worth re-examining. Otherwise we've kind of locked ourselves in to only looking at dose reconstruction and -- and -- and -- and not the site profile, and it may be the very fertile, raw material out of which each of the cookie-cutter dose reconstructions in some cases are going to be extracted. So I just would encourage you all to rethink whether it's maybe time to have a recommendation to NIOSH in a formal way urging some

kind of conflict of interest provisions which mirror those that are used by Dr. Toohey and others at Oak Ridge Associated Universities on the dose reconstruction conflicts. I just -- I don't know how many more of these you want to have sprout up before you finally take Board recommendation in this area. 7 I would just offer this with respect to the evening meeting question. I think it varies. I think it goes from 8 site to site. If you've got well-organized claimants or well-organized institutions that can help you do it, you know, it's where you are. Right? I mean I think there'll be lots of volunteers in western New York and, you know, maybe you'll have them, maybe you won't have 3 them. And I think it ought to be -- I think you ought to figure out in advance -- if you're going to schedule an evening meeting, figure out if it's going to be productive -- right? -- and -- and not just do it and then have an empty room and sit there for two hours and close the record. So -- and I'm certainly happy to help NIOSH network with people where they're not already plugged in. And I'm sure there are others on 2 and off the Board that would be delighted to do so

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because I think it would be immensely productive and it
      would also promote greater understanding. I think
      there needs to be a two-way communication on this
      'cause it's complicated.
5 Finally, I would just propose -- probably for the last time
      I'll raise it publicly, here at least -- but I -- I
      have -- will -- will sort of -- I guess raise the
      question -- I'm delighted that the Board is going to be
      -- apparently going to get trained and learn to use
      IMBA and all of those good things and -- and -- and I
      was delighted to read that if you want to go to
      Cincinnati, you can, even as a member of the public,
      have an opportunity to use IMBA. I -- I think there
      need to be some creative solutions to dealing with the
      proprietary software issue. I don't know how much
      creativity's been applied to it at this point, or maybe
      there's a way your support service con-- you know, Oak
      Ridge, could provide some kind of service so that if
      you don't want to provide proprietary software, at
      least you can make somebody available who can, for
      members of the public, make use of IMBA and IMBA
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      outputs. It's -- it's a very difficult thing to take a
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blank site profile which are where the doses are conveyed and -- particularly the internals are conveyed and, you know, DPMs or whatever with solubilities identified and -- and be able to replicate the results that NIOSH is providing to individual claimants. And -- but more particularly, it becomes more important as you get into things like extrapolation where you don't have data and -- and I -- I would just encourage you all to think about whether it makes sense to have a program relying on proprietary dose reconstruction software where it effectively is inaccessible to the public except in the most -- except with a very high barrier, meaning that -- that the price tag for interacting with NIOSH or with this program or even this Board at a technical level is ponying up some six figures or five or six figures to get access to software. And -- and I -- I know I -- I respect that, you know, people are in the business of making money on their software programs, but you've got a public program here -- public compensation program relying on private software. And it -- it -- I think it poses a transparency question and I would hope that you

the third time; as I say, it will be the last time I revisit it here -- but I -- I -- I just think you've got a problem assuring transparency throughout, and this is a huge obstacle in that transparency, so I would encourage some thought to that. Thank you. Thankful -- or thank you, Richard, for your 7 DR. ZIEMER: thoughtful comments. Let's take a minute and see if Board members have questions to pose to you relative to your remarks. Any? 1 DR. MELIUS: Yeah, I have one question and I -- for both I guess Larry and Richard, and that's the Blockson 2 Chemical -- if I understand that correctly -- and I 3 don't understand it completely, but what's an issue -more of a generic issue with a number of the AEC sites in particular where there's a sort of a commercial use and a -- commercial exposures or exposures from other industrial process as well as from the AEC process, and the issue is how you parse the exposures and do them, and I -- and I don't understand to what extent it's policy, legal or whatever, but I think certainly a 2 briefing on that would -- at some point soon would be

all would -- once again, I'm revisiting this now for

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very helpful to the Board so at least we understand
      what -- what's involved in the decision-making, as well
      as how it will affect future dose reconstructions,
      so...
5 DR. ZIEMER: And Richard also entered into the public record
      some related comments. I think the Board members have
      gotten copies of your written comments on that Biloxi
      issue, Richard.
9 Let me ask a somewhat related question. Does this revolve
      around the official definition of -- of the facility
      insofar as it relates to weapons production?
2 MR. MILLER: You know what, it's like lawyers say, it
      depends who you ask.
4 DR. ZIEMER: Well, but -- but that -- that lies at the crux
      of it, as the starting point that it's defined in a
      certain way and then that gets interpreted. Well,
      where is that line exactly? Does it cover this -- the
      -- clearly there's radon related to the phosphate
      thing. I know -- I see Dr. Roessler here, who worked
      with phosphates a lot down in Florida and very much
      aware of the radon issues, but then the issue is where
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      -- where does that end as far as what that company was
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doing anyway and where does the -- the uranium work
      begin and were the uranium workers also exposed and is
      this part of their occupational exposure.
                                                 That's --
                It's a great set of questions, Dr. Ziemer,
4 MR. MILLER:
      and it's --
6 DR. ZIEMER: Yeah, I --
7 MR. MILLER: Well, go ahead. I mean -- I didn't mean to cut
      you off.
             Well, and I think Jim is saying we -- we don't
9 DR. ZIEMER:
      necessarily know what all those issues are, either.
      understand that perhaps NIOSH has itself been
      addressing that or looking at that, and I don't know if
      you want to make any comments on that, but that's
3
      certainly been part of the issue. You're -- you're
      perhaps questioning whether or not the decision has
      been made and is fixed in concrete, and it may affect
                                  Is that --
      other facilities, as well.
              Well, I think there's several things.
8 MR. MILLER:
      -- I don't know what the final status of the
      interagency deliberations are in this. I know there's
      certainly some options and comments that have been
      circulated. I guess the question is, let's leave aside
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the interagency debates for a moment and just step back by bringing to the fore what actually this Board brings, which is a remarkable rich diversity of expertise, and ask the question -- assuming that an atomic weapons employer facility lawfully encompasses a facility, broadly defined, and you're looking at the production of materials that are ultimately used for a nuclear weapons program where you have this dual-use issue -- right? -- commercial and non -- and non -military and non-military, we'll say, or DOE and non-DOE -- where you then have to ask the question where it may even be inseparable, and let's assume that legally you can look at the whole thing. And then let's say okay, where does it make sense to tease it out? other words, can we apply common sense to this question, 'cause it's a thought puzzle, I think. It's a thought puzzle. I mean -- you know, having sat down and kind of sketched out about five options, I could persuasively -- to myself, at least -- argue five different ways to draw the line on this thing. But in fact it's a -- partly a health physics question and it's partly a engineering question and it's partly a

policy call about how you deal with the equities for individuals, and it is -- I mean -- and -- and one could just -- I mean the myriad of equity issues just jump out at you. Right? Well, the -- so the person who makes rock phosphate and gets radon exp-- you know, processes this into phosphoric acid and they have radon exposures but they make Tide and they're not compensable, but the individuals who -- who make -- who -- who are part of the -- how do you even parse it out because the same person's making it for both -- both production chains. I don't have the -- the ri-- in other words, I'm not here advocating a particular solution, but I do think the options ought to be fleshed out because it's -- it is so important to the equities about whether or not you attribute dose that is at lea-- least partially attributable to the work that you're doing that wound up as part of this company's work for the Atomic Energy Commission or not. And it's not so remote that it's laughable. Right? It's just -- it's an equity issue. It's -- It's a Solomon-like activity, where are you going to divide the baby? And I think reasonable people could differ

on this thing, but I think it ought to be aired out and -- and I think it ought not be decided simply in an interagency deliberation process. Or to the extent it is, at least it would be useful if the perspectives of this Board were also informing the thinking of the Federal -- Federally-responsible officials in this respect. That -- that's sort of my pitch.

8 DR. ZIEMER: Thank you.

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9 MR. MILLER: Let me -- let me just -- just add one last thing, which is -- is -- is since poor Larry often seems like my -- my good friend Eeyore who -- who -- who dreads coming to these meetings and -- and looks very unhappy most of the time, and I want to say something nice for Larry for a change, before I say something else, and -- so you can get off your pins and needles now. I found -- NIOSH's work on the site profile at Mallinckrodt begs numerous questions. Denise raised terrific questions, and I have them, as well. How are you going to deal with the periods of time where you don't have good dose information or any dose information, and how can we have any confidence that we're not estimating -- underestimating the

periods where you don't have dose data, and particularly when you're relying on, you know, 50-yearold methods of analysis. But in the course of the site profile, I found a terrific footnote for a document that at least I had been looking for for over a year, diligently, with numerous requests all over the place, by Merril Eisenbud and had -- was very pleased to see that it was footnoted in the site profile, and -- and in this Eisenbud document -- I don't know if the Board has had a chance to look at it, but it was the basis for an article that ran in the Riverfront Times, it's -- it's called an estimate of cumulative multiple exposures to radioactive materials at Mallinckrodt's plants four and six from November of 1950. What was stunning, in addition to the level of doses that were estimated for a typical worker in the matrix that was used, which was about 1,000 rem to the lung over about a two-and-a-half-year period, was -- was the -- and this formerly secret memo said, and it just -- just to tickle your fancy, it -- it -- if I could just read one paragraph to you, it says here (reading) Early in 1947 the New York operations office evaluated the potential

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hazards in these plants and, after finding it to be considerable, recommended the necessary corrective actions. In addition, steps were taken by NYOO in cooperation with the contractor to institute procedures for effective environmental and personal monitoring. It was recognized that pending elimination of excessive exposures, here was a unique opportunity to conduct clinical studies on a fairly large-sized population whose radiation exposure for several years had been considerably in excess of any group for which data was available.

2 And I just have to say that this really was a remarkable window in a candid memo about the perspectives that were in place at least at that time about how convenient it was to study the workers at Mallinckrodt without regard to how they were put in harm's way. And that really was the foundation of a lot of what informed the passage of this law. And every now and then these -- these -- these remarkable historical documents finally see their way to light and -- and I just would like to thank NIOSH for making sure that this one found its way to light and -- and they

- produced it without need for a FOIA request and did so
- in a transparent way. And so I just wanted to express
- 3 that appreciation.
- 4 DR. ZIEMER: Thank you. Thank you, Larry -- or thank you,
- Richard. Thank you both.
- 6 Okay, that concludes our public comment requests.
- 7 MS. BROCK: (Off microphone) Dr. Ziemer --
- 8 MR. ELLIOTT: She's thought of something.
- 9 MS. BROCK: -- (Off microphone) can I make one more comment?
- O DR. ZIEMER: Yes, you may. Denise Brock.
- 1 MS. BROCK: I forgot to mention earlier, I -- in reference
- to what Cori and -- and Larry, I believe, were talking
- about how to alert people or media. I actually had a
- 4 list recently that I used that actually listed all of
- the papers -- suburban journals, any paper that I could
- 6 come up with through the state of Missouri and through
- 17 Illinois, as well as the -- the news channels and the
- 8 radios, and I sent letters out to each and every one of
- them actually looking for any of the 3,300 employees of
- Mallinckrodt or anybody involved in the building and
- 1 construction trades. And I've gotten a lot of response
- \$2 from these papers saying that they are going to run ads

looking for -- for people that worked at these facilities. 3 I was also contacted by some people in Illinois who are wanting me to come there to help them organize and kind of do what we did for the Mallinckrodt site, so I'm going to do that, as well. 7 But I was curious if it's ever possible to do like a public service announcement, if that's possible, to put something like that on TV and maybe just try to get contact people in each area to alert claimants or to try to -- and I think that's probably part of the outreach with the Department of Labor, as well. Thank you. That's very helpful. We then are 3 DR. ZIEMER: at the end of our open business session. Let me ask if any of the Board members have any additional items or comments for the good of the order? 'Cause if they don't, we are going to adjourn the public session. will take our lunch break a little early. I think we probably -- well, in fact they do need to clear this room out anyway and prepare it for the big show tonight, so you need to get all of your stuff out of 2 here.

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the Mesquite I Room at 2:00 o'clock in the Mesquite I Room --
the Mesquite I Room at 2:00 o'clock. I will, for the
record, again emphasize that this is a session that is
only for the discussion of the task orders and the
independent government cost estimate. The Board will
do no other business at that meeting.
We now stand adjourned as far as the public meeting is
concerned. Thank you very much.
(Whereupon, the public portion of the meeting was
adjourned.)
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2 Error!

CERTIFICATE

Error!
STATE OF GEORGIA)
COUNTY OF FULTON)

I, STEVEN RAY GREEN, being a Certified Merit Court
Reporter in and for the State of Georgia, do hereby certify
that the foregoing transcript was reduced to typewriting by
me personally or under my direct supervision, and is a true,
complete, and correct transcript of the aforesaid
proceedings reported by me.

I further certify that I am not related to, employed by, counsel to, or attorney for any parties, attorneys, or counsel involved herein; nor am I financially interested in this matter.

WITNESS MY HAND AND OFFICIAL SEAL this _____ day of January, 2004.

S	STEVEN		RAY	GREEN,	CVR-CM
G2	Α	CCR	No.	A-2102	